

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: March 9, 2005, 16:53:39 ; Search time 156 Seconds
(without alignments)
230.757 Million cell updates/sec

Title: US-10-613-228A-1

Perfect score: 22

Sequence: 1 tcgcgttttcgtcgttttc 22

Scoring table:

IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 1202784 seqs, 818138359 residues

Total number of hits satisfying chosen parameters: 2405568

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :

Issued Patents NA: *
1: /cgn2_6/ptodata/1/ina/5A_COMB.seq: *
2: /cgn2_6/ptodata/1/ina/5B_COMB.seq: *
3: /cgn2_6/ptodata/1/ina/6A_COMB.seq: *
4: /cgn2_6/ptodata/1/ina/6B_COMB.seq: *
5: /cgn2_6/ptodata/1/ina/PCCTS_COMB.seq: *
6: /cgn2_6/ptodata/1/ina/backfiles1.seq: *

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	17.2	78.2	133	US-09-313-294A-108	Sequence 108, App
2	17.2	78.2	513	US-09-107-532A-292	Sequence 292, App
3	16.8	76.4	36016	US-09-949-016-14223	Sequence 14223, A
4	16.4	74.5	3531	US-08-629-600-1	Sequence 1, Appl
5	16.4	74.5	3531	US-09-076-761-1	Sequence 1, Appl
6	16.2	73.6	909	US-09-134-000C-3167	Sequence 3167, App
7	16.2	73.6	2358	US-09-134-000C-3285	Sequence 3285, App
8	15.8	71.8	660	US-09-489-039A-560	Sequence 560, App
9	15.6	70.9	231	US-09-543-681A-2772	Sequence 2772, App
10	15.6	70.9	300	US-09-543-681A-1478	Sequence 1478, App
11	15.6	70.9	440	US-09-513-999C-35185	Sequence 35185, A
12	15.6	70.9	599	US-09-270-767-3552	Sequence 3552, App
13	15.6	70.9	599	US-09-270-767-18834	Sequence 18834, A
14	15.6	70.9	601	US-09-949-016-31352	Sequence 31352, A
15	15.6	70.9	601	US-09-949-016-69221	Sequence 69221, A
16	15.6	70.9	601	US-09-949-016-81370	Sequence 81370, A
17	15.6	70.9	601	US-09-949-016-127420	Sequence 127420, A
18	15.6	70.9	601	US-09-949-016-171125	Sequence 171125, A
19	15.6	70.9	601	US-09-949-016-171126	Sequence 171126, A
20	15.6	70.9	601	US-09-949-016-171127	Sequence 171127, A
21	15.6	70.9	628	US-09-134-001C-2551	Sequence 2551, App
22	15.6	70.9	826	US-09-221-017B-1008	Sequence 1008, App
23	15.6	70.9	850	US-08-617-860B-34	Sequence 34, Appl
24	15.6	70.9	1068	US-09-543-681A-2972	Sequence 2972, App
25	15.6	70.9	1734	US-09-248-796A-4704	Sequence 4704, App
26	15.6	70.9	2109	US-09-248-796A-2999	Sequence 2999, App
27	15.6	70.9	2394	US-09-540-236-893	Sequence 893, App

28	15.6	70.9	3069	US-08-335-865J-7	Sequence 7, Appl
29	15.6	70.9	3805	US-09-513-729B-10	Sequence 10, Appl
30	15.6	70.9	3805	US-09-023-655-1443	Sequence 1443, App
31	15.6	70.9	4071	US-09-513-057C-5	Sequence 5, Appl
32	15.6	70.9	4071	US-09-746-801A-5	Sequence 5, Appl
33	15.6	70.9	4285	US-09-949-016-689	Sequence 689, App
34	15.6	70.9	4308	US-09-394-142B-23	Sequence 23, Appl
35	15.6	70.9	4643	US-08-605-106-6	Sequence 6, Appl
36	15.6	70.9	5061	US-09-355-160D-1	Sequence 1, Appl
37	15.6	70.9	5061	US-10-092-219-1	Sequence 1, Appl
38	15.6	70.9	5296	US-09-949-016-2362	Sequence 2362, App
39	15.6	70.9	8302	US-09-234-827B-1	Sequence 1, Appl
40	15.6	70.9	10204	US-09-949-016-14104	Sequence 14104, A
41	15.6	70.9	10482	US-09-322-478-23	Sequence 23, Appl
42	15.6	70.9	10482	US-09-586-106D-23	Sequence 23, Appl
43	15.6	70.9	14066	US-09-601-198-56	Sequence 56, Appl
44	15.6	70.9	26709	US-09-949-016-17520	Sequence 17520, A
45	15.6	70.9	34279	US-09-596-002-26	Sequence 26, Appl

ALIGNMENTS

```
RESULT 1
US-09-313-294A-108
; Sequence 108, Application US/09313294A
; Patent No. 6476212
; GENERAL INFORMATION:
; APPLICANT: Ialagudi, Raghunath V.
; APPLICANT: Ito, Laura Y.
; APPLICANT: Sherman, Bradley K.
; TITLE OF INVENTION: POLYNUCLEOTIDES AND POLYPEPTIDES DERIVED FROM CORN EAR
; FILE REFERENCE: PU-0017 US
; CURRENT FILING DATE: US/09/313, 294A
; NUMBER OF SEQ ID NOS: 1999-05-14
; SOFTWARE: PERL Program
; SEQ ID NO 108
; LENGTH: 133
; TYPE: DNA
; ORGANISM: Zea mays
; FEATURE:
; NAME/KEY: misc feature
; OTHER INFORMATION: Incyte ID No. 6476212 700548545H1
; NAME/KEY: unsure
; LOCATION: 8
; OTHER INFORMATION: a, t, c, g, or other
US-09-313-294A-108

Query Match      78.2% Score 17.2; DB 4; Length 133;
Best Local Similarity 86.4%; Pred. No. 1.2e+02;
Matches 19; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Cy      1 TCgcgttttcgcgttttc 22
Db      72 TCgcgttttcgcgttttc 93

RESULT 2
US-09-107-532A-292/C
; Sequence 292, Application US/09107532A
; Patent No. 6583275
; GENERAL INFORMATION:
; APPLICANT: Lynn A Doucette-Stamm and David Bush
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO
; ENTEROCOCCUS FACIUM FOR DIAGNOSTICS AND THERAPEUTICS
; NUMBER OF SEQUENCES: 7310
; CORRESPONDENCE ADDRESS:
; ADDRESS: GENOME THERAPEUTICS CORPORATION
; STREET: 100 Beaver Street
; CITY: Waltham
; STATE: Massachusetts
; COUNTRY: USA
```

```

1      ZIP: 02354
2      COMPUTER READABLE FORM:
3      MEDIUM TYPE: CD/ROM ISO9660
4      COMPUTER: PC
5      OPERATING SYSTEM: <Unknown>
6      SOFTWARE: ASCII
7      CURRENT APPLICATION DATA:
8      APPLICATION NUMBER: US/09/107,532A
9      FILING DATE: 30-Jun-1998
10     PRIOR APPLICATION DATA:
11     APPLICATION NUMBER: 60/085,598
12     FILING DATE: 14 May 1998
13     APPLICATION NUMBER: 60/051571
14     FILING DATE: July 2, 1997
15     ATTORNEY/AGENT INFORMATION:
16     NAME: Ariadello, Pamela Deneke
17     REGISTRATION NUMBER: 40,489
18     REFERENCE/DOCKET NUMBER: GTC-012
19     TELECOMMUNICATION INFORMATION:
20     TELEPHONE: (781)893-5007
21     TELEFAX: (781)893-8277
22     INFORMATION FOR SEQ ID NO: 292:
23     SEQUENCE CHARACTERISTICS:
24     LENGTH: 513 base pairs
25     TYPE: nucleic acid
26     STRANDEDNESS: double
27     TOPOLOGY: circular
28     MOLECULE TYPE: DNA (genomic)
29     HYPOTHETICAL: NO
30     ANTI-SENSE: NO
31     ORIGINAL SOURCE:
32     ORGANISM: Enterococcus faecium
33     FEATURE:
34     NAME/KEY: misc feature
35     LOCATION: (B) LOCATION 1...513
36     SEQUENCE DESCRIPTION: SEQ ID NO: 292:
37     US-09-107-532A-292

```

Query Match	78.2%	Score 17.2	DB 4	Length 513
Best Local Similarity	86.4%	Pred. No. 1.3e+02		
Matches 19	Conservative 0	Mismatches 3	Indels 0	Gaps 0

[illegible]

```

RESULT 3
US-09-949-016-14223/c
; Sequence 14223, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CLO01107
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 14223
; LENGTH: 36016
; TYPE: DNA
; ORGANISM: Human
; US-09-949-016-14223

```

Query Match	76.4%; Score 16.8; DB 4; Length 36016;
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```

Qy      3 GTCGTTTTTCGTGGCTTTT 22
          |||||
Db    31422 GTCGTTTTTTTGTCGTTTTT 31403
Best Local Similarity 90.0%; Pred. No. 2,7e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
```

RESULT 4
 US-08-629-600-1/c
 Sequence 1, Application US/08629600
 Patent No. 5783196
 GENERAL INFORMATION:
 APPLICANT: NORIEGA, Fernando
 APPLICANT: LEVINE, Myron M.
 TITLE OF INVENTION: GUA MUTANTS OF SHIGELLA
 TITLE OF INVENTION: AND VACCINES CONTAINING THE SAME
 NUMBER OF SEQUENCES: 18
 CORRESPONDENCE ADDRESSES:
 ADDRESSSEE: SUGHRU, MION, ZINN, MACPEAK & SEAS
 STREET: 2100 Pennsylvania Avenue, N.W., Suite 800
 CITY: Washington, D.C.
 STATE: D.C.
 COUNTRY: U.S.A.
 ZIP: 20037
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: Patentin Release #1.0, Version #1.25
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/629,600
 FILING DATE: 9-Apr-1996
 CLASSIFICATION: 424
 ATTORNEY/AGENT INFORMATION:
 NAME: KIT, Gordon
 REGISTRATION NUMBER: 30,764
 REFERENCE/DOCKET NUMBER: A-6765
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (202) 293-7060
 TELEFAX: (202) 293-7860
 INFORMATION FOR SEQ ID NO: 1:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 3531 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 MOLECULE TYPE: genomic DNA
 HYPOTHETICAL: NO
 ANTI-SENSE: NO
 US-08-629-600-1

Query Match	74.5%	Score 16.4	DB 1	Length 3531
Best Local Similarity	94.4%	Pred. No. 3.3e+02		
Matches 17	Conservative	0	Mismatches 1	Indels 0
Gaps				0

QY	5	CGTTTTCGTCGCTTTT	22
Db	846	CGTTTTCGTCGCTTTT	829

```

RESULT 5
US-09-076-761-1/c
: Sequence 1, Application US/09076761
: Patent No. 6190669
:
: GENERAL INFORMATION:
:
: APPLICANT: NORIEGA, Fernando
: APPLICANT: SZTEIN, Marcelo B.
: APPLICANT: LEVINE, Myron M.
: TITLE OF INVENTION: ATTENUATED MUTANTS OF SALMONELLA
: TITLE OF INVENTION: WHICH CONSTITUTIVELY EXPRESS THE
: TITLE OF INVENTION: Vi ANTIGEN
: NUMBER OF SEQUENCES: 10

```

CORRESPONDENCE ADDRESS:

ADDRESSEE: SUGHRUE, MION, ZINN, MACPEAK & SEAS
STREET: 2100 Pennsylvania Avenue, N.W., Suite 800
CITY: Washington, D.C.
STATE: D.C.
COUNTRY: U.S.A.
ZIP: 20037

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/076,761
FILING DATE: 13-MAY-1998
CLASSIFICATION:
ATTORNEY/AGENT INFORMATION:

NAME: KIT, Gordon
REGISTRATION NUMBER: 30,764
REFERENCE/DOCKET NUMBER: A-7140
TELECOMMUNICATION INFORMATION:
TELEPHONE: (202) 293-7060
TELEFAX: (202) 293-7860
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 3531 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: genomic DNA
HYPOTHETICAL: NO
ANTI-SENSE: NO
US-09-076-761-1

Query Match 74.5%; Score 16.4; DB 3; Length 3531;
Best Local Similarity 94.4%; Pred. No. 3.3e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 5 CGTTTTCGTGCGTTT 22
DB 846 CGTTTTCGTGCGATT 829

RESULT 6

US-09-134-000C-3167/C
Sequence 3167, Application US/09134000C
Patent No. 6617156
GENERAL INFORMATION:
APPLICANT: Lynn Doucette-Stamm et al
TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO
FILE REFERENCE: 032796-032
CURRENT APPLICATION NUMBER: US/09/134,000C
CURRENT FILING DATE: 1998-08-13
PRIOR APPLICATION NUMBER: US 60/055,778
PRIOR FILING DATE: 1997-08-15
NUMBER OF SEQ ID NOS: 6812
SOFTWARE: Patentin version 3.1
SEQ ID NO 3167
LENGTH: 909
TYPE: DNA
ORGANISM: Enterococcus faecalis
US-09-134-000C-3167

Query Match 73.6%; Score 16.2; DB 4; Length 909;
Best Local Similarity 85.7%; Pred. No. 3.7e+02;
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 CGTCGTTTTCGTGCGTTT 22
DB 248 CGTCGTTTTCGTGCGTTGT 228

RESULT 7

US-09-134-000C-3285/C
Sequence 3285, Application US/09134000C
Patent No. 6617156
GENERAL INFORMATION:
APPLICANT: Lynn Doucette-Stamm et al
TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO
FILE REFERENCE: 032796-032
CURRENT APPLICATION NUMBER: US/09/134,000C
CURRENT FILING DATE: 1998-08-13
PRIOR APPLICATION NUMBER: US 60/055,778
PRIOR FILING DATE: 1997-08-15
NUMBER OF SEQ ID NOS: 6812
SOFTWARE: Patentin version 3.1
SEQ ID NO 3285
LENGTH: 2358
TYPE: DNA
ORGANISM: Enterococcus faecalis
US-09-134-000C-3285

Query Match 73.6%; Score 16.2; DB 4; Length 2358;
Best Local Similarity 85.7%; Pred. No. 4e+02;
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 CGTCGTTTTCGTGCGTTT 22
DB 1697 CGTCGTTTTCGTGCGTTGT 1677

RESULT 8

US-09-489-039A-560
Sequence 560, Application US/09489039A
Patent No. 6610836
GENERAL INFORMATION:
APPLICANT: Gary Breton et. al
TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO KLEBSIELLA
FILE REFERENCE: 2709 2004001
CURRENT APPLICATION NUMBER: US/09/489,039A
CURRENT FILING DATE: 2000-01-27
PRIOR APPLICATION NUMBER: US 60/117,747
PRIOR FILING DATE: 1999-01-29
NUMBER OF SEQ ID NOS: 14342
SEQ ID NO 560
LENGTH: 660
TYPE: DNA
ORGANISM: Klebsiella pneumoniae
US-09-489-039A-560

Query Match 71.8%; Score 15.8; DB 4; Length 660;
Best Local Similarity 89.5%; Pred. No. 5.4e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 3 GTCGTTTTCGTGCGTTT 21
DB 426 GTCGTTTTCGTGCGATT 444

RESULT 9

US-09-543-681A-2772
Sequence 2772, Application US/09543681A
Patent No. 6605709
GENERAL INFORMATION:
APPLICANT: GARY BRETON
TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PROTEUS MIRABILIS
FILE REFERENCE: 2709 1002-001
CURRENT APPLICATION NUMBER: US/09/543,681A
CURRENT FILING DATE: 2000-04-05
PRIOR APPLICATION NUMBER: US 60/128,706
PRIOR FILING DATE: 1999-04-09
NUMBER OF SEQ ID NOS: 8344

```
; SEQ ID NO 2772
;
; LENGTH: 231
;
; TYPE: DNA
; ORGANISM: Proteus mirabilis
US-09-543-681A-2772
```

Query Match	70.9%;	Score 15.6;	DB 4;	Length 231;
Best Local Similarity	81.8%;	Pred. No. 6.1e+02;		
Matches 18; Conservative	0;	Mismatches 4;	Indels 0;	Gaps 0

Qy		1 TCCTCGTTTTCGTGCGTTTTT	22
Db		167 TCCTCGTTTTCAGCTTTTTT	188

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RESULT 10
US-09-543-681A-1478/c
; Sequence 1478, Application US/09543681A
; Patent No. 6605709
; GENERAL INFORMATION:
; APPLICANT: GARY BRETON
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PROTEUS MIRABILIS
; TITLE OF INVENTION: DIAGNOSTICS AND THERAPEUTICS
; FILE REFERENCE: 2709.1002-001
; CURRENT APPLICATION NUMBER: US/09/543,681A
; CURRENT FILING DATE: 2000-04-05
; PRIOR APPLICATION NUMBER: US 60/128,706
; PRIOR FILING DATE: 1999-04-09
; NUMBER OF SEQ ID NOS: 8344
; SEQ ID NO 1478
; LENGTH: 300
; TYPE: DNA
; ORGANISM: Proteus mirabilis
; US-09-543-681A-1478

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Query Match	70.9%;	Score 15.6;	DB 4;	Length 300;
Best Local Similarity	81.8%;	Pred. No. 6.2e+02;		
Matches 18; Conservative	0;	Mismatches 4;	Indels 0;	Gaps 0;

```

QY      1 TCGTCGTTTTTCGTGCGGTTTT 22
          ||||| ||||| ||
Db      208 TCGTGGTTTTTCTGCGGTGTT 187

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```

RESULT 11
US-09-513-999C-35185
; Sequence 35185, Application US/09513999C
; Patent No. 6783961
; GENERAL INFORMATION:
; APPLICANT: Dumas Milne Edwards, J. B.
; APPLICANT: Duclert, A.
; APPLICANT: Giordano, J. Y.
; TITLE OF INVENTION: Expressed Sequence Tags and Encoded Human Proteins
; Patent No. 6783961
; FILE REFERENCE: 59.US2.REG
; CURRENT APPLICATION NUMBER: US/09/513,999C
; CURRENT FILING DATE: 2000-02-24
; PRIOR APPLICATION NUMBER: US 60/122,487
; PRIOR FILING DATE: 1999-02-26
; NUMBER OF SEQ ID NOS: 36681
; SOFTWARE: Patent.pm
; SEQ ID NO 35185
; LENGTH: 440
; TYPE: DNA
; ORGANISM: Homo sapiens
; US-09-513-999C-35185

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Query Match	70.9%;	Score 15.6;	DB 4;	Length 440;
Best Local Similarity	81.8%;	Pred. No. 6.4e+02;		
Matches 18; Conservative	0;	Mismatches 4;	Indels 0;	Gaps 0;

QY 1 TCGTCGTTTTCTCGTCGTTTTT 22
| | | | | | | | | | | | | |

Db 193 TTGTCCTTTTTTTTGCGTTTTT 214

```

RESULT 12
US-09-270-767-3552/c
; Sequence 3552, Application US/09270767
; Patent No. 6703481
; GENERAL INFORMATION:
; APPLICANT: Homburger et al.
; TITLE OF INVENTION: Nucleic acids and proteins of Drosophila melanogaster
; FILE REFERENCE: File Reference: 7326-094
; CURRENT APPLICATION NUMBER: US/09/270,767
; CURRENT FILING DATE: 1999-03-17
; NUMBER OF SEQ ID NOS: 62517
; SOFTWARE: Patencin Ver. 2.0
; SEQ ID NO 3552
; LENGTH: 599
; TYPE: DNA
; ORGANISM: Drosophila melanogaster
US-09-270-767-3552

```

Query Match	70.9%;	Score 15.6;	DB 4;	Length 599;
Best Local Similarity	81.8%;	Pred. No. 6.5e+02;		
Matches 18; Conservative	0;	Mismatches 4;	Indels 0;	Gaps 0;

QY		1	TGTCGTTTTTCGTGCCTTTT	22
Db		568	TCGTCGTGTCGTGTTTTTTT	547

```

RESULT 13
US-09-270-767-18834/c
; Sequence 18834, Application US/09270767
; Patent No. 6703421
; GENERAL INFORMATION:
; APPLICANT: Homburger et al.
; TITLE OF INVENTION: Nucleic acids and proteins of Drosophila melanogaster
; FILE REFERENCE: File Reference: 7326-094
; CURRENT APPLICATION NUMBER: US/09/270,767
; CURRENT FILING DATE: 1999-03-17
; NUMBER OF SEQ ID NOS: 62517
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 18834
; LENGTH: 599
; TYPE: DNA
; ORGANISM: Drosophila melanogaster
US-09-270-767-18834

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Query Match	70.9%;	Score 15.6;	DB 4;	Length 599;
Best Local Similarity	81.8%;	Pred. No. 6.5e+02;		
Matches 18; Conservative	0;	Mismatches 4;	Indels 0;	Gaps 0;

QY	1	TCGTCGTTTTTCGTGCGTTTTT	22
Db	568	TCGTCGTGTTTCGTGTTTTTTT	547

RESULT 14
 US-09-949-016--31352/C
 ; Sequence 31352, Application US/09949016
 ; Patent No. 6812339
 ;
 ; GENERAL INFORMATION:
 ;
 ; APPLICANT: VENTER, J. Craig et al.
 ;
 ; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
 ; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
 ;
 ; FILE REFERENCE: C1001307
 ;
 ; CURRENT APPLICATION NUMBER: US/09/949,016
 ;
 ; CURRENT FILING DATE: 2000-04-14
 ;
 ; PRIOR APPLICATION NUMBER: 60/241,755
 ;
 ; PRIOR FILING DATE: 2000-10-20
 ;
 ; PRIOR APPLICATION NUMBER: 60/237,768
 ;
 ; PRIOR FILING DATE: 2000-10-03
 ;
 ; PRIOR APPLICATION NUMBER: 60/231,498

;
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 31352
; LENGTH: 601
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-31352

Query Match 70.9%; Score 15.6; DB 4; Length 601;
Best Local Similarity 81.8%; Pred. No. 6.5e+02;
Matches 18; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

Qy 1 TCGTCGTTTTTCGTCGCGTTTTT 22
 ||||| ||||| |||||
Db 334 TTGTCCTTTTTCGTCGTTTTT 313

RESULT 15
US-09-949-016-69221/C
; Sequence 69221. Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE REFERENCE: C1001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 69221
; LENGTH: 601
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-69221

Query Match 70.9%; Score 15.6; DB 4; Length 601;
Best Local Similarity 81.8%; Pred. No. 6.5e+02;
Matches 18; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

Qy 1 TCGTCGTTTTTCGTCGCGTTTTT 22
 ||||| ||||| |||||
Db 416 TCTTCGTTTTCTCGGCGTTTTT 395

Search completed: March 9, 2005, 18:48:14
Job time : 158 secs

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OM nucleic - nucleic search, using sw model

Run on: March 9, 2005, 18:45:43 ; Search time 578 Seconds
(without alignments)
225.856 Million cell updates/sec

Title: US-10-613-228A-1

Perfect score: 22

Sequence: 1 tcgcgttttcgcgttttc 22

Scoring table:

IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 5401638 seqs, 2966923429 residues

Total number of hits satisfying chosen parameters: 10803276

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Published Applications NA:*

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16: /cgn2_6/ptodata/1/pubpna/US10D_PUBCOMB.seq:*
17: /cgn2_6/ptodata/1/pubpna/US10E_PUBCOMB.seq:*
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21: /cgn2_6/ptodata/1/pubpna/US60_NEW_PUB.seq:*
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	22	100.0	22	17	US-10-613-228A-1
2	22	100.0	22	18	US-10-816-220-152
3	20.4	92.7	618	18	US-10-363-345A-34257
4	20.4	92.7	618	18	US-10-363-345A-34258
5	20.4	92.7	619	18	US-10-363-345A-40287
6	20.4	92.7	619	18	US-10-363-345A-40288
7	20.4	92.7	761	18	US-10-363-345A-2025
8	20.4	92.7	761	18	US-10-363-345A-2026
9	20.4	92.7	1024	18	US-10-363-345A-7204
10	20.4	92.7	1024	18	US-10-363-345A-7204
11	19.4	88.2	511	18	US-10-363-345A-28015

C 12	19.4	88.2	511	18	US-10-363-345A-28016	Sequence 28016, A
C 13	19.4	88.2	523	18	US-10-363-345A-35059	Sequence 35059, A
C 14	19.4	88.2	523	18	US-10-363-345A-35060	Sequence 35060, A
C 15	19.4	88.2	524	18	US-10-363-345A-40325	Sequence 40325, A
C 16	19.4	88.2	524	18	US-10-363-345A-40326	Sequence 40326, A
C 17	19.4	88.2	610	18	US-10-363-345A-23153	Sequence 23153, A
C 18	19.4	88.2	610	18	US-10-363-345A-23154	Sequence 23154, A
C 19	19.4	88.2	838	18	US-10-363-345A-32131	Sequence 32131, A
C 20	19.4	88.2	838	18	US-10-363-345A-32132	Sequence 32132, A
C 21	19.4	88.2	839	18	US-10-363-345A-1741	Sequence 1741, Ap
C 22	19.4	88.2	839	18	US-10-363-345A-1742	Sequence 1742, Ap
C 23	19.4	88.2	885	18	US-10-363-345A-18089	Sequence 18089, A
C 24	19.4	88.2	885	18	US-10-363-345A-18090	Sequence 18090, A
C 25	19.4	88.2	1267	18	US-10-363-345A-27261	Sequence 27261, A
C 26	19.4	88.2	1267	18	US-10-363-345A-27262	Sequence 27262, A
C 27	19.4	88.2	3673778	16	US-10-312-841-2	Sequence 2, Appl
C 28	19	86.4	920	18	US-10-363-345A-20285	Sequence 20285, A
C 29	19	86.4	920	18	US-10-363-345A-20286	Sequence 20286, A
C 30	18.8	85.5	523	18	US-10-363-345A-17495	Sequence 17495, A
C 31	18.8	85.5	523	18	US-10-363-345A-17496	Sequence 17496, A
C 32	18.8	85.5	524	18	US-10-363-345A-15539	Sequence 15539, A
C 33	18.8	85.5	524	18	US-10-363-345A-15540	Sequence 15540, A
C 34	18.8	85.5	525	18	US-10-363-345A-35507	Sequence 35507, A
C 35	18.8	85.5	525	18	US-10-363-345A-35508	Sequence 35508, A
C 36	18.8	85.5	553	18	US-10-363-345A-39041	Sequence 39041, A
C 37	18.8	85.5	553	18	US-10-363-345A-39042	Sequence 39042, A
C 38	18.8	85.5	561	18	US-10-363-345A-35971	Sequence 35971, A
C 39	18.8	85.5	561	18	US-10-363-345A-35972	Sequence 35972, A
C 40	18.8	85.5	651	18	US-10-363-345A-34913	Sequence 34913, A
C 41	18.8	85.5	651	18	US-10-363-345A-34914	Sequence 34914, A
C 42	18.8	85.5	712	18	US-10-363-345A-27451	Sequence 27451, A
C 43	18.8	85.5	712	18	US-10-363-345A-27452	Sequence 27452, A
C 44	18.8	85.5	759	18	US-10-363-345A-3263	Sequence 3263, Ap
C 45	18.8	85.5	759	18	US-10-363-345A-3264	Sequence 3264, Ap

ALIGNMENTS

RESULT 1
US-10-613-228A-1
; Sequence 1, Application US/10613228A
; Publication No. US20040092472A1
; GENERAL INFORMATION:
; APPLICANT: KRIEG, ARTHUR M
; TITLE OF INVENTION: NUCLEIC ACID COMPOSITIONS FOR STIMULATING IMMUNE RESPONSES
; FILE REFERENCE: C1037, 70045080
; CURRENT APPLICATION NUMBER: US/10/613, 228A
; CURRENT FILING DATE: 2003-07-03
; PRIOR APPLICATION NUMBER: US 60/394,193
; PRIOR FILING DATE: 2002-07-03
; NUMBER OF SEQ ID NOS: 3
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 1
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial sequence
; FEATURE:
; OTHER INFORMATION: Oligodeoxynucleotide
US-10-613-228A-1

Query Match 100.0%; Score 22; DB 17; Length 22;
Best Local Similarity 100.0%; Pred. No. 3.8;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCgcgttttcgcgttttc 22
|||
Db 1 TCgcgttttcgcgttttc 22

RESULT 2
US-10-816-220-152
; Sequence 152, Application US/10816220

```
; Publication No. US20040235770A1
; GENERAL INFORMATION:
; APPLICANT: DAVIS, Heather L
; APPLICANT: McCLUSKIE, Michael J
; TITLE OF INVENTION: IMMUNOSTIMULATORY NUCLEIC ACID OIL-IN-WATER FORMULATIONS AND
; TITLE OF INVENTION: RELATED METHODS OF USE
; FILE REFERENCE: C1037.70039US01
; CURRENT APPLICATION NUMBER: US/10/816,220
; CURRENT FILING DATE: 2004-04-01
; PRIOR FILING DATE: 2003-04-02
; PRIOR APPLICATION NUMBER: US 60/461,903
; PRIOR FILING DATE: 2003-04-10
; NUMBER OF SEQ ID NOS: 434
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 152
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Oligonucleotide
US-10-816-220-152

Query Match          100.0%; Score 22; DB 18; Length 22;
Best Local Similarity 100.0%; Pred. No. 3.8;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCGTCGTTTTTCGTCGTTTTT 22
Db 1 TCGTCGTTTTTCGTCGTTTTT 22

RESULT 3
US-10-363-345A-34257
; Sequence 34257, Application US/10363345A
; Publication No. US20040234960A1
; GENERAL INFORMATION:
; APPLICANT: Alexander Olek
; APPLICANT: Christian Piepenbrock
; APPLICANT: Kurt Berlin
; TITLE OF INVENTION: Method for determining the degree of methylation of defined
; TITLE OF INVENTION: cytosines in genomic DNA in the sequence context of 5'-CpG-3
; FILE REFERENCE: E01/1227
; CURRENT APPLICATION NUMBER: US/10/363,345A
; CURRENT FILING DATE: 2003-03-03
; NUMBER OF SEQ ID NOS: 40712
; SEQ ID NO 34257
; LENGTH: 618
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)
; OTHER INFORMATION: CpG-island No: 34257
US-10-363-345A-34257

Query Match          92.7%; Score 20.4; DB 18; Length 618;
Best Local Similarity 95.5%; Pred. No. 27;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TCGTCGTTTTTCGTCGTTTTT 22
Db 98 TCGTCGTTTTTCGTCGTTTTT 119

RESULT 4
US-10-363-345A-34258/c
; Sequence 34258, Application US/10363345A
; Publication No. US20040234960A1
; GENERAL INFORMATION:
; APPLICANT: Alexander Olek
; APPLICANT: Christian Piepenbrock
; APPLICANT: Kurt Berlin
; TITLE OF INVENTION: Method for determining the degree of methylation of defined
```

```
; TITLE OF INVENTION: cytosines in genomic DNA in the sequence context of 5'-CpG-3
; FILE REFERENCE: E01/1227
; CURRENT APPLICATION NUMBER: US/10/363,345A
; CURRENT FILING DATE: 2003-03-03
; NUMBER OF SEQ ID NOS: 40712
; SEQ ID NO 34258
; LENGTH: 618
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)
; OTHER INFORMATION: CpG-island No: 34258
US-10-363-345A-34258

Query Match          92.7%; Score 20.4; DB 18; Length 618;
Best Local Similarity 95.5%; Pred. No. 27;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TCGTCGTTTTTCGTCGTTTTT 22
Db 521 TCGTCGTTTTTCGTCGTTTTT 500

RESULT 5
US-10-363-345A-40287
; Sequence 40287, Application US/10363345A
; Publication No. US20040234960A1
; GENERAL INFORMATION:
; APPLICANT: Alexander Olek
; APPLICANT: Christian Piepenbrock
; APPLICANT: Kurt Berlin
; TITLE OF INVENTION: Method for determining the degree of methylation of defined
; TITLE OF INVENTION: cytosines in genomic DNA in the sequence context of 5'-CpG-3
; FILE REFERENCE: E01/1227
; CURRENT APPLICATION NUMBER: US/10/363,345A
; CURRENT FILING DATE: 2003-03-03
; NUMBER OF SEQ ID NOS: 40712
; SEQ ID NO 40287
; LENGTH: 619
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)
; OTHER INFORMATION: CpG-island No: 40287
US-10-363-345A-40287

Query Match          92.7%; Score 20.4; DB 18; Length 619;
Best Local Similarity 95.5%; Pred. No. 27;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TCGTCGTTTTTCGTCGTTTTT 22
Db 190 TCGTCGTTTTTCGTCGTTTTT 211

RESULT 6
US-10-363-345A-40288/c
; Sequence 40288, Application US/10363345A
; Publication No. US20040234960A1
; GENERAL INFORMATION:
; APPLICANT: Alexander Olek
; APPLICANT: Christian Piepenbrock
; APPLICANT: Kurt Berlin
; TITLE OF INVENTION: Method for determining the degree of methylation of defined
; TITLE OF INVENTION: cytosines in genomic DNA in the sequence context of 5'-CpG-3
; FILE REFERENCE: E01/1227
; CURRENT APPLICATION NUMBER: US/10/363,345A
; CURRENT FILING DATE: 2003-03-03
; NUMBER OF SEQ ID NOS: 40712
; SEQ ID NO 40288
; LENGTH: 619
; TYPE: DNA
; ORGANISM: Artificial Sequence
```



```
; FEATURE:
; OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)
; OTHER INFORMATION: Cpg-Island No: 40288
US-10-363-345A-40288
```

```
Query Match          92.7%; Score 20.4; DB 18; Length 619;
Best Local Similarity 95.5%; Pred. No. 27;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
```

```
Qy 1 TCGTCGTTTTCGTCGCGTTTTT 22
Db 430 TCGTCGTTTTCGTCGCGTTTTT 409
```

RESULT 7

```
US-10-363-345A-2025
; Sequence 2025, Application US/10363345A
; Publication No. US20040234960A1
; GENERAL INFORMATION:
; APPLICANT: Alexander Olek
; APPLICANT: Christian Piepenbrock
; APPLICANT: Kurt Berlin
; TITLE OF INVENTION: Method for determining the degree of methylation of defined
; FILE REFERENCE: E01/1227
; CURRENT APPLICATION NUMBER: US/10/363,345A
; CURRENT FILING DATE: 2003-03-03
; NUMBER OF SEQ ID NOS: 40712
; SEQ ID NO 2025
; LENGTH: 761
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)
; OTHER INFORMATION: Cpg-Island No: 2025
US-10-363-345A-2025
```

```
Query Match          92.7%; Score 20.4; DB 18; Length 761;
Best Local Similarity 95.5%; Pred. No. 28;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
```

```
Qy 1 TCGTCGTTTTCGTCGCGTTTTT 22
Db 598 TCGTCGTTTTCGTCGCGTTTTT 619
```

RESULT 8

```
US-10-363-345A-2026/c
; Sequence 2026, Application US/10363345A
; Publication No. US20040234960A1
; GENERAL INFORMATION:
; APPLICANT: Alexander Olek
; APPLICANT: Christian Piepenbrock
; APPLICANT: Kurt Berlin
; TITLE OF INVENTION: Method for determining the degree of methylation of defined
; FILE REFERENCE: E01/1227
; CURRENT APPLICATION NUMBER: US/10/363,345A
; CURRENT FILING DATE: 2003-03-03
; NUMBER OF SEQ ID NOS: 40712
; SEQ ID NO 2026
; LENGTH: 761
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)
; OTHER INFORMATION: Cpg-Island No: 2026
US-10-363-345A-2026
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```
Query Match          92.7%; Score 20.4; DB 18; Length 761;
Best Local Similarity 95.5%; Pred. No. 28;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
```

```
Qy 1 TCGTCGTTTTCGTCGCGTTTTT 22
Db 164 TCGTCGTTTTCGTCGCGTTTTT 143
```

RESULT 9

```
US-10-363-345A-7203
; Sequence 7203, Application US/10363345A
; Publication No. US20040234960A1
; GENERAL INFORMATION:
; APPLICANT: Alexander Olek
; APPLICANT: Christian Piepenbrock
; APPLICANT: Kurt Berlin
; TITLE OF INVENTION: Method for determining the degree of methylation of defined
; FILE REFERENCE: E01/1227
; CURRENT APPLICATION NUMBER: US/10/363,345A
; CURRENT FILING DATE: 2003-03-03
; NUMBER OF SEQ ID NOS: 40712
; SEQ ID NO 7203
; LENGTH: 1024
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)
; OTHER INFORMATION: Cpg-Island No: 7203
US-10-363-345A-7203
```

```
Query Match          92.7%; Score 20.4; DB 18; Length 1024;
Best Local Similarity 95.5%; Pred. No. 29;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
```

```
Qy 1 TCGTCGTTTTCGTCGCGTTTTT 22
Db 276 TCGTCGTTTTCGTCGCGTTTTT 297
```

RESULT 10

```
US-10-363-345A-7204/c
; Sequence 7204, Application US/10363345A
; Publication No. US20040234960A1
; GENERAL INFORMATION:
; APPLICANT: Alexander Olek
; APPLICANT: Christian Piepenbrock
; APPLICANT: Kurt Berlin
; TITLE OF INVENTION: Method for determining the degree of methylation of defined
; FILE REFERENCE: E01/1227
; CURRENT APPLICATION NUMBER: US/10/363,345A
; CURRENT FILING DATE: 2003-03-03
; NUMBER OF SEQ ID NOS: 40712
; SEQ ID NO 7204
; LENGTH: 1024
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)
; OTHER INFORMATION: Cpg-Island No: 7204
US-10-363-345A-7204
```

```
Query Match          92.7%; Score 20.4; DB 18; Length 1024;
Best Local Similarity 95.5%; Pred. No. 29;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
```

```
Qy 1 TCGTCGTTTTCGTCGCGTTTTT 22
Db 749 TCGTCGTTTTCGTCGCGTTTTT 728
```

RESULT 11

```
US-10-363-345A-28015
; Sequence 28015, Application US/10363345A
; Publication No. US20040234960A1
```

```
; GENERAL INFORMATION:
; APPLICANT: Alexander Olek
; APPLICANT: Christian Piepenbrock
; APPLICANT: Kurt Berlin
; TITLE OF INVENTION: Method for determining the degree of methylation of defined
; FILE REFERENCE: E01/1227
; CURRENT APPLICATION NUMBER: US/10/363,345A
; CURRENT FILING DATE: 2003-03-03
; NUMBER OF SEQ ID NOS: 40712
; SEQ ID NO 28015
; LENGTH: 511
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)
; OTHER INFORMATION: Cpg-island No: 28015
US-10-363-345A-28015
```

```
Query Match      88.2%; Score 19.4; DB 18; Length 511;
Best Local Similarity 95.2%; Pred. No. 72;
Matches 20; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
```

```
QY      2 CGTCGTTTTCGTCGCGTTTTT 22
Db      474 CGTCGTTTTCGGCGCGTTTTT 494
```

```
RESULT 12
US-10-363-345A-28016/c
; Sequence 28016, Application US/10363345A
; Publication No. US20040234960A1
; GENERAL INFORMATION:
; APPLICANT: Alexander Olek
; APPLICANT: Christian Piepenbrock
; APPLICANT: Kurt Berlin
; TITLE OF INVENTION: Method for determining the degree of methylation of defined
; FILE REFERENCE: E01/1227
; CURRENT APPLICATION NUMBER: US/10/363,345A
; CURRENT FILING DATE: 2003-03-03
; NUMBER OF SEQ ID NOS: 40712
; SEQ ID NO 28016
; LENGTH: 511
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)
; OTHER INFORMATION: Cpg-island No: 28016
US-10-363-345A-28016
```

```
Query Match      88.2%; Score 19.4; DB 18; Length 511;
Best Local Similarity 95.2%; Pred. No. 72;
Matches 20; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
```

```
QY      2 CGTCGTTTTCGTCGCGTTTTT 22
Db      38 CGTCGTTTTCGGCGCGTTTTT 18
```

```
RESULT 13
US-10-363-345A-35059
; Sequence 35059, Application US/10363345A
; Publication No. US20040234960A1
; GENERAL INFORMATION:
; APPLICANT: Alexander Olek
; APPLICANT: Christian Piepenbrock
; APPLICANT: Kurt Berlin
; TITLE OF INVENTION: Method for determining the degree of methylation of defined
; FILE REFERENCE: E01/1227
; CURRENT APPLICATION NUMBER: US/10/363,345A
; CURRENT FILING DATE: 2003-03-03
```

```
; NUMBER OF SEQ ID NOS: 40712
; SEQ ID NO 35059
; LENGTH: 523
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)
; OTHER INFORMATION: Cpg-island No: 35059
US-10-363-345A-35059
```

```
Query Match      88.2%; Score 19.4; DB 18; Length 523;
Best Local Similarity 95.2%; Pred. No. 73;
Matches 20; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
```

```
QY      2 CGTCGTTTTCGTCGCGTTTTT 22
Db      488 CGTCGTTTTCGTCGCGTTTTT 508
```

```
RESULT 14
US-10-363-345A-35060/c
; Sequence 35060, Application US/10363345A
; Publication No. US20040234960A1
; GENERAL INFORMATION:
; APPLICANT: Alexander Olek
; APPLICANT: Christian Piepenbrock
; APPLICANT: Kurt Berlin
; TITLE OF INVENTION: Method for determining the degree of methylation of defined
; FILE REFERENCE: E01/1227
; CURRENT APPLICATION NUMBER: US/10/363,345A
; CURRENT FILING DATE: 2003-03-03
; NUMBER OF SEQ ID NOS: 40712
; SEQ ID NO 35060
; LENGTH: 523
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)
; OTHER INFORMATION: Cpg-island No: 35060
US-10-363-345A-35060
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```
Query Match      88.2%; Score 19.4; DB 18; Length 523;
Best Local Similarity 95.2%; Pred. No. 73;
Matches 20; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
```

```
QY      2 CGTCGTTTTCGTCGCGTTTTT 22
Db      36 CGTCGTTTTCGTCGCGTTTTT 16
```

```
RESULT 15
US-10-363-345A-40325
; Sequence 40325, Application US/10363345A
; Publication No. US20040234960A1
; GENERAL INFORMATION:
; APPLICANT: Alexander Olek
; APPLICANT: Christian Piepenbrock
; APPLICANT: Kurt Berlin
; TITLE OF INVENTION: Method for determining the degree of methylation of defined
; FILE REFERENCE: E01/1227
; CURRENT APPLICATION NUMBER: US/10/363,345A
; CURRENT FILING DATE: 2003-03-03
; NUMBER OF SEQ ID NOS: 40712
; SEQ ID NO 40325
; LENGTH: 524
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)
; OTHER INFORMATION: Cpg-island No: 40325
US-10-363-345A-40325
```

Query Match 88.2%; Score 19.4; DB 18; Length 524;
 Best Local Similarity 95.2%; Pred. No. 73;
 Matches 20; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 2 CGTCGTTTTTCGTGCGTTTTT 22
 |||||
 Db 399 CGTCGTTTTTCGTGCGTTTTT 419

Search completed: March 9, 2005, 21:05:42
 Job time : 581 secs

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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: March 9, 2005, 14:14:33 ; Search time 3888 Seconds
(without alignments)
274.181 Million cell updates/sec

Title: US-10-613-228A-1

Perfect score: 22
Sequence: 1 tcgtcgcttttcgtcgcttttc 22

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 4708233 seqs, 24227607955 residues

Total number of hits satisfying chosen parameters: 9416466

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%

Listing first 45 summaries

Database :

GenEmbl:*
1: gb_ba:*
2: gb_hcg:*
3: gb_in:*
4: gb_om:*
5: gb_ov:*
6: gb_pat:*
7: gb_ph:*
8: gb_pl:*
9: gb_pr:*
10: gb_ro:*
11: gb_sts:*
12: gb_sy:*
13: gb_un:*
14: gb_vt:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	20.4	92.7	226153	2 AC073704	Mus muscu
2	19.4	88.2	349980	6 AX344573	Sequence
3	18.8	85.5	5826	6 AX356381	Sequence
4	18.8	85.5	6130	6 AX251400	Sequence
5	18.8	85.5	6130	6 AX345031	Sequence
6	18.8	85.5	7728	6 AX344979	Sequence
7	18.8	85.5	7728	6 AX348500	Sequence
8	18.8	85.5	8951	6 AX345697	Sequence
9	18.8	85.5	13133	6 AX344227	Sequence
10	18.4	83.6	20	6 AX104184	Sequence
11	18.4	83.6	20	6 AX355698	Sequence
12	18.4	83.6	20	6 AX547237	Sequence
13	18.4	83.6	5369	6 CO600212	Sequence
14	18.4	83.6	146491	2 AC019950	Drosophi
15	18.4	83.6	182601	3 AC009904	Drosophi
16	18.4	83.6	310675	3 AE003708	Drosophi
17	18.4	83.6	310675	1 AE016869	Pseudom
18	18.4	83.6	149269	2 BX897667	Danio rer
19	17.8	80.9	1527	5 BC056691	Danio rer

20	17.8	80.9	2290	3 AF606934	Auf606934 Spirula s
21	17.8	80.9	2322	8 AY139685	AY139685 Porphyra
22	17.8	80.9	2381	6 AX347239	AX347239 Sequence
23	17.8	80.9	7195	6 AX277866	AX277866 Sequence
24	17.8	80.9	7195	6 AX323551	AX323551 Sequence
25	17.8	80.9	11622	6 AX345576	AX345576 Sequence
26	17.8	80.9	34216	3 U57054	U57054 Caenorhabdi
27	17.8	80.9	99176	5 BX255893	BX255893 Zebrafish
28	17.8	80.9	172585	2 CR847782	CR847782 Danio rer
29	17.8	80.9	215917	2 AC006764	AC006764 Caenorhab
30	17.8	80.9	252250	1 AP005339	AP005339 Vibrio vu
31	17.8	80.9	300045	1 AE016803	AE016803 Vibrio vu
32	17.8	80.9	303121	1 AE016766	AE016766 Escherich
33	17.4	79.1	4110	6 AX598855	AX598855 Sequence
34	17.4	79.1	5774	6 AX278031	AX278031 Sequence
35	17.4	79.1	5774	6 AX323798	AX323798 Sequence
36	17.4	79.1	5938	6 AX344811	AX344811 Sequence
37	17.4	79.1	6067	6 AX344680	AX344680 Sequence
38	17.4	79.1	6124	8 AB010408	AB010408 Arabidops
39	17.4	79.1	7004	6 AX277956	AX277956 Sequence
40	17.4	79.1	7004	6 AX323651	AX323651 Sequence
41	17.4	79.1	7110	6 CQ806871	CQ806871 Sequence
42	17.4	79.1	7110	6 AX251243	AX251243 Sequence
43	17.4	79.1	7110	6 AX251990	AX251990 Sequence
44	17.4	79.1	7110	6 AX346458	AX346458 Sequence
45	17.4	79.1	7110	6 AX349019	AX349019 Sequence

ALIGNMENTS

RESULT 1
AC073704
LOCUS
DEFINITION Mus musculus clone RP23-175112, WORKING DRAFT SEQUENCE, 35
unordered pieces.
AC073704
AC073704.1 GI:8810321
HTG; HTGS_PHASE1; HTGS_DRAFT.
KEYWORDS Mus musculus (house mouse)
SOURCE
ORGANISM
REFERENCE
AUTHORS
TITLE
JOURNAL
REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT

Center: Joint Genome Institute
Center Code: JGI
Web site: <http://www.jgi.doe.gov>

Project Information
Center Project Name: 1804276
Center clone name: RPCI-23_175112

Summary Statistics
Consensus quality: 200795 bases at least Q40
Consensus quality: 211634 bases at least Q30
Consensus quality: 213977 bases at least Q20
Estimated insert size: 205410; agarose-ff estimation
Estimated insert size: 222753; sum-of-contigs estimation
Quality coverage: 7.81 in Q20 bases; agarose-ff estimation
Quality coverage: 7.2 in Q20 bases; sum-of-contigs estimation.
* NOTE: This is a 'working draft' sequence. It currently
* consists of 35 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as

* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence.
* as soon as it is available and the accession number will
* be preserved.
* 1
* 1252: contig of 1252 bp in length
* 1253
* 1352: gap of unknown length
* 1353
* 2189: contig of 1037 bp in length
* 2390
* 2489: gap of unknown length
* 2490
* 4017: contig of 1528 bp in length
* 4018
* 4117: gap of unknown length
* 4118
* 5570: contig of 1153 bp in length
* 5271
* 5370: gap of unknown length
* 5371
* 6704: contig of 1334 bp in length
* 6705
* 6805
* 6805
* 7662: contig of 1158 bp in length
* 7963
* 8063
* 9083: contig of 1021 bp in length
* 9084
* 9184
* 10470: contig of 1287 bp in length
* 10471
* 10570: gap of unknown length
* 10571
* 11849: contig of 1279 bp in length
* 11850
* 11949: gap of unknown length
* 11950
* 13311: contig of 1362 bp in length
* 13312
* 13311: gap of unknown length
* 13412
* 14471: contig of 1060 bp in length
* 14472
* 14571: gap of unknown length
* 14572
* 15881: contig of 1310 bp in length
* 15882
* 15981: gap of unknown length
* 15982
* 17316: contig of 1335 bp in length
* 17317
* 17316: gap of unknown length
* 17417
* 19718: contig of 1802 bp in length
* 19719
* 19318: gap of unknown length
* 19319
* 20320: contig of 1002 bp in length
* 20321
* 20420: gap of unknown length
* 20421
* 21903: contig of 1483 bp in length
* 21904
* 22003: gap of unknown length
* 22004
* 23728: contig of 1725 bp in length
* 23729
* 23828: gap of unknown length
* 23829
* 26109: contig of 2281 bp in length
* 26110
* 26210
* 27815: contig of 1606 bp in length
* 27816
* 27915: gap of unknown length
* 27916
* 30044: contig of 2129 bp in length
* 30045
* 30144: gap of unknown length
* 30145
* 35020: contig of 4876 bp in length
* 35021
* 35120: gap of unknown length
* 35121
* 40883: contig of 5763 bp in length
* 40884
* 40883: gap of unknown length
* 40984
* 46087: contig of 5104 bp in length
* 46088
* 46187: gap of unknown length
* 46188
* 51176: contig of 5529 bp in length
* 51177
* 51816: gap of unknown length
* 51817
* 59638: contig of 7822 bp in length
* 59639
* 59738: gap of unknown length
* 59739
* 66829: contig of 7091 bp in length
* 66830
* 66929: gap of unknown length
* 66930
* 78269: contig of 11340 bp in length
* 78270
* 78369: gap of unknown length
* 78370
* 88909: contig of 10540 bp in length
* 88910
* 89010
* 101128: gap of unknown length
* 101129
* 101228: gap of unknown length
* 101229
* 116533: contig of 15225 bp in length
* 11654
* 116533: gap of unknown length
* 116554
* 130907: contig of 14354 bp in length
* 130908
* 131007: gap of unknown length
* 131008
* 146644: contig of 15857 bp in length
* 146685
* 146664: gap of unknown length
* 146965
* 171782: contig of 24818 bp in length
* 171783
* 171882: gap of unknown length
* 171883
* 198608: contig of 26262 bp in length
* 198609
* 198608: gap of unknown length
* 198609
* 226153: contig of 27545 bp in length.

FEATURES	location/Qualifiers
SOURCE	1. .226153
	/organism="Mus musculus"
	/mol_type="genomic DNA"
	/db_xref="taxon:10090"
	/clone="RP23-175112"
	/clone_1lb="RPC1 mouse BAC library 23"
ORIGIN	
Query Match	92.7%; Score 20.4; DB 2; Length 226153;
Best Local Similarity	95.5%; Pred. No. 87;
Matches	21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
OY	1 TCGTCGTTTTTCGTCGCTTTT 22
Db	20672 TCGTCGTTTTTCGTCGCTTTT 20693
RESULT 2	
LOCUS	AX344573 349980 bp DNA linear PAT 01-FEB-2002
DEFINITION	Sequence 24 from Patent WO0200932.
ACCESSION	AX344573
VERSION	AX344573.1 GI:18492459
KEYWORDS	
SOURCE	synthetic construct
ORGANISM	synthetic construct
REFERENCE	1
AUTHORS	Olek A., Piepenbrock C. and Berlin K.
TITLE	Diagnosis of known genetic disorders within the mhc
JOURNAL	Patent: WO 0200932-A 24 03-JAN-2002;
FEATURES	Epigenomics AG (DE)
SOURCE	location/Qualifiers
	1. .349980
	/organism="synthetic construct"
	/mol_type="unassigned DNA"
	/db_xref="taxon:32630"
	/note="chemically treated genomic DNA (Homo sapiens).-Original length of seq 1: 3.673778 <223>-split as follows.-seq 01 0.000.001 TO 0.349.980-seq 02 0.300.001 649.980-seq 03 600.001 949.980-seq 04 900.001 1.249.980-seq 05 1.200.001 1.549.980-seq 06 1.500.001 1.449.980-seq 07 1.800.001 2.149.980-seq 08 2.100.001 2.449.980-seq 09 2.400.001 2.749.980-seq 10 2.700.001 3.049.980-seq 11 3.000.001 3.349.980-seq 12 3.300.001 3.649.980-seq 13 3.600.001 3.673.778 <223>-Original length of seq 2: 3.673778 <223>-split as follows.-seq 14 0.000.001 TO 0.349.980-seq 15 0.300.001 649.980-seq 16 600.001 949.980-seq 17 900.001 1.249.980-seq 18 1.200.001 1.549.980-seq 19 1.500.001 1.849.980-seq 20 1.800.001 2.149.980-seq 21 2.100.001 2.449.980-seq 22 2.400.001 2.749.980-seq 23 2.700.001 3.049.980-seq 24 3.000.001 3.349.980-seq 25 3.300.001 3.649.980-seq 26 3.600.001 3.673.778"
ORIGIN	
Query Match	88.2%; Score 19.4; DB 6; Length 349980;
Best Local Similarity	95.2%; Pred. No. 2,3e+02;
Matches	20; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
OY	1 TCGTCGTTTTTCGTCGCTTTT 21
Db	213728 TCGTCGTTTTTCGTCGCTTTT 213748
RESULT 3	
LOCUS	AX356381 5826 bp DNA linear PAT 06-FEB-2002
DEFINITION	Sequence 15 from Patent WO0181622.
ACCESSION	AX356381
VERSION	AX356381.1 GI:18620871
KEYWORDS	

SOURCE synthetic construct
ORGANISM synthetic construct
other sequences; artificial sequences.

REFERENCE 1
AUTHORS Olek, A., Piepenbrock, C. and Berlin, K.
TITLE Diagnosis of diseases associated with dna repair
JOURNAL Patent: WO 0181622-A 15 01-NOV-2001;
Epigenomics AG (DE)

FEATURES
source location/Qualifiers
1..5826
/organism="synthetic construct"
/mol_type="unassigned DNA"
/db_xref="taxon:32630"
/note="Chemically treated genomic DNA (Homo sapiens)"

ORIGIN

Query Match 85.5%; Score 18.8; DB 6; Length 5826;
Best Local Similarity 90.9%; Pred. No. 6.4e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TCGTCGTTTTCGTCGCTTTT 22
|||||
Db 5567 TCGTCGTTTTCGTCGCTTTT 5588

RESULT 4
LOCUS AX251400 6130 bp DNA linear PAT 05-OCT-2001
DEFINITION Sequence 368 from Patent WO0168912.
ACCESSION AX251400
VERSION AX251400.1 GI:15984823
KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
other sequences; artificial sequences.

REFERENCE 1
AUTHORS Olek, A., Piepenbrock, C. and Berlin, K.
TITLE Diagnosis of diseases associated with tumor suppressor genes and oncogenes
JOURNAL Patent: WO 0168912-A 368 20-SEP-2001;
Epigenomics AG (DE)

FEATURES
source location/Qualifiers
1..6130
/organism="synthetic construct"
/mol_type="unassigned DNA"
/db_xref="taxon:32630"
/note="Chemically treated genomic DNA (Homo sapiens)"

ORIGIN

Query Match 85.5%; Score 18.8; DB 6; Length 6130;
Best Local Similarity 90.9%; Pred. No. 6.4e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TCGTCGTTTTCGTCGCTTTT 22
|||||
Db 1332 TCGGCCTTTTCGTCGCTTTT 1353

RESULT 5
LOCUS AX345031 6130 bp DNA linear PAT 01-FEB-2002
DEFINITION Sequence 102 from Patent WO0200928.
ACCESSION AX345031
VERSION AX345031.1 GI:18492917
KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
other sequences; artificial sequences.

REFERENCE 1
AUTHORS Olek, A., Piepenbrock, C. and Berlin, K.
TITLE Diagnosis of diseases associated with the immune system
JOURNAL Patent: WO 0200928-A 102 03-JAN-2002;
Epigenomics AG (DE)

FEATURES
source location/Qualifiers
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/organism="synthetic construct"
/mol_type="unassigned DNA"
/db_xref="taxon:32630"
/note="Chemically treated genomic DNA (Homo sapiens)"

ORIGIN

Query Match 85.5%; Score 18.8; DB 6; Length 6130;
Best Local Similarity 90.9%; Pred. No. 6.4e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TCGTCGTTTTCGTCGCTTTT 22
|||||
Db 1332 TCGGCCTTTTCGTCGCTTTT 1353

RESULT 6
LOCUS AX344979 7728 bp DNA linear PAT 01-FEB-2002
DEFINITION Sequence 50 from Patent WO0200928.
ACCESSION AX344979
VERSION AX344979.1 GI:18492865
KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
other sequences; artificial sequences.

REFERENCE 1
AUTHORS Olek, A., Piepenbrock, C. and Berlin, K.
TITLE Diagnosis of diseases associated with the immune system
JOURNAL Patent: WO 0200928-A 50 03-JAN-2002;
Epigenomics AG (DE)

FEATURES
source location/Qualifiers
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/organism="synthetic construct"
/mol_type="unassigned DNA"
/db_xref="taxon:32630"
/note="Chemically treated genomic DNA (Homo sapiens)"

ORIGIN

Query Match 85.5%; Score 18.8; DB 6; Length 7728;
Best Local Similarity 90.9%; Pred. No. 6.3e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TCGTCGTTTTCGTCGCTTTT 22
|||||
Db 2026 TCGTCGTTTTCGTCGCTTTT 2047

RESULT 7
LOCUS AX348500 7728 bp DNA linear PAT 06-FEB-2002
DEFINITION Sequence 8 from Patent WO0202809.
ACCESSION AX348500
VERSION AX348500.1 GI:18614535
KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
other sequences; artificial sequences.

REFERENCE 1
AUTHORS Olek, A., Piepenbrock, C. and Berlin, K.
TITLE Diagnosis of behavioural disorders, neurological disorders and Cancer
JOURNAL Patent: WO 0202809-A 8 10-JAN-2002;
Epigenomics AG (DE)

FEATURES
source location/Qualifiers
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/organism="synthetic construct"
/mol_type="unassigned DNA"
/db_xref="taxon:32630"
/note="Chemically treated genomic DNA (Homo sapiens)"

ORIGIN

Query Match 85.5%; Score 18.8; DB 6; Length 7728;
Best Local Similarity 90.9%; Pred. No. 6.3e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 1 TCGTCGTTTTCGTCGCTTTT 22
|||||
Db 2026 TCGTCGTTTTCGTCGCTTTT 2047

RESULT 8
AX345697 8951 bp DNA linear PAT 01-FEB-2002
LOCUS Sequence 768 from Patent WO0200928.
DEFINITION AX345697
ACCESSION AX345697
VERSION AX345697.1 GI:18493583
KEYWORDS
SOURCE synthetic construct
ORGANISM other sequences; artificial sequences.

REFERENCE 1
AUTHORS Olek, A., Piepenbrock, C. and Berlin, K.
TITLE Diagnosis of diseases associated with the immune system
JOURNAL Patent: WO 0200928-A 768 03-JAN-2002;
Epigenomics AG (DE)

FEATURES
source Location/Qualifiers
1. 8951
/organism="synthetic construct"
/mol_type="unassigned DNA"
/db_xref="taxon:32630"
/note="chemically treated genomic DNA (Homo sapiens)"

ORIGIN

Query Match 85.5%; Score 18.8; DB 6; Length 8951;
Best Local Similarity 90.9%; Pred. No. 6.2e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TCGTCGTTTTCGTCGCTTTT 22
|||||
Db 3724 TCGCGCTTTTCGTCGCTTTT 3745

RESULT 9
AX344227 13133 bp DNA linear PAT 01-FEB-2002
LOCUS AX344227
DEFINITION Sequence 74 from Patent WO0200926.
ACCESSION AX344227
VERSION AX344227.1 GI:18492115
KEYWORDS
SOURCE synthetic construct
ORGANISM other sequences; artificial sequences.

REFERENCE 1
AUTHORS Olek, A., Piepenbrock, C. and Berlin, K.
TITLE Diagnosis of diseases associated with signal transduction
JOURNAL Patent: WO 0200926-A 74 03-JAN-2002;
Epigenomics AG (DE)

FEATURES
source Location/Qualifiers
1. 13133
/organism="synthetic construct"
/mol_type="unassigned DNA"
/db_xref="taxon:32630"
/note="chemically treated genomic DNA (Homo sapiens)"

ORIGIN

Query Match 85.5%; Score 18.8; DB 6; Length 13133;
Best Local Similarity 90.9%; Pred. No. 6e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TCGTCGTTTTCGTCGCTTTT 22
|||||
Db 8050 TCGTCGTTTTCGTCGCTTTT 8071

RESULT 10
AX104184 20 bp DNA linear PAT 30-APR-2001
LOCUS AX104184
DEFINITION Sequence 376 from Patent WO0122972.
ACCESSION AX104184
VERSION AX104184.1 GI:13920381
KEYWORDS
SOURCE synthetic construct
ORGANISM other sequences; artificial sequences.

REFERENCE 1
AUTHORS Krieg, A.M., Schetter, C. and Vollmer, J.C.
TITLE Immunostimulatory nucleic acids
JOURNAL Patent: WO 0122972-A 376 05-APR-2001;
UNIVERSITY OF IOWA RESEARCH FOUNDATION (US) ; Coley Pharmaceutical
GmbH (DE)

FEATURES
source Location/Qualifiers
1. 20
/organism="synthetic construct"
/mol_type="unassigned DNA"
/db_xref="taxon:32630"

ORIGIN

Query Match 83.6%; Score 18.4; DB 6; Length 20;
Best Local Similarity 95.0%; Pred. No. 1.7e+03;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TCGTCGTTTTCGTCGCTTT 20
|||||
Db 1 TCGTCGTTTTCGTCGCTTT 20

RESULT 11
AX355698 20 bp DNA linear PAT 06-FEB-2002
LOCUS AX355698
DEFINITION Sequence 726 from Patent WO0197843.
ACCESSION AX355698
VERSION AX355698.1 GI:18620366
KEYWORDS
SOURCE synthetic construct
ORGANISM other sequences; artificial sequences.

REFERENCE 1
AUTHORS Weiner, G. and Hartmann, G.
TITLE Methods for enhancing antibody-induced cell lysis and treating
JOURNAL Patent: WO 0197843-A 726 27-DEC-2001;
UNIVERSITY OF IOWA RESEARCH FOUNDATION (US)

FEATURES
source Location/Qualifiers
1. 20
/organism="synthetic construct"
/mol_type="unassigned DNA"
/db_xref="taxon:32630"
/note="Synthetic oligonucleotide-phosphorothioate
backbone"

ORIGIN

Query Match 83.6%; Score 18.4; DB 6; Length 20;
Best Local Similarity 95.0%; Pred. No. 1.7e+03;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TCGTCGTTTTCGTCGCTTT 20
|||||
Db 1 TCGTCGTTTTCGTCGCTTT 20

RESULT 12
AX547237 20 bp DNA linear PAT 01-MAR-2003
LOCUS AX547237
DEFINITION Sequence 376 from Patent WO02053141.
ACCESSION AX547237
VERSION AX547237.1 GI:25812381
KEYWORDS

SOURCE synthetic construct
ORGANISM synthetic construct
other sequences; artificial sequences.

REFERENCE 1 Bratzler, R.L.
TITLE Inhibition of angiogenesis by nucleic acids
JOURNAL Patent: WO 02053141-A 376 11-JUL-2002;
Coley Pharmaceutical Group, Inc. (US)

FEATURES
source
1. 20
/organism="synthetic construct"
/mol_type="unassigned DNA"
/db_xref="taxon:32630"
/note="Synthetic Sequence"

ORIGIN

Query Match 83.6%; Score 18.4; DB 6; Length 20;
Best Local Similarity 95.0%; Pred. No. 1.7e+03;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TCGTCGTTTTGTCGCGTTT 20
1 TCGTCGTTTTGTCGCGTTT 20

Db 1 TCGTCGTTTTGTCGCGTTT 20

RESULT 13
LOCUS CQ600212 5369 bp DNA linear PAT 02-FEB-2004
DEFINITION Sequence 27970 from Patent WO0171042.
ACCESSION CQ600212
VERSION CQ600212.1 GI:41655454
KEYWORDS
SOURCE Drosophila sp.
ORGANISM Drosophila sp.
Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
Ephydroidea; Drosophilidae; Drosophila.

REFERENCE 1 Venter, J.C., Adams, M., Li, P.W. and Myers, E.W.
TITLE Detection kites, such as nucleic acid arrays, for detecting the
JOURNAL expression of 10,000 or more Drosophila genes and uses thereof
Patent: WO 0171042-A 27970 27-SEP-2001;
PE Corporation (NY) (US)

FEATURES
source
1. 5369
/organism="Drosophila sp."
/mol_type="unassigned DNA"
/db_xref="taxon:7242"

ORIGIN

Query Match 83.6%; Score 18.4; DB 6; Length 5369;
Best Local Similarity 95.0%; Pred. No. 9.8e+02;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 CGTCGTTTTGTCGCGTTT 21
3568 CGTCGTTTTGTCGCGTTT 3587

Db 3568 CGTCGTTTTGTCGCGTTT 3587

RESULT 14
LOCUS AC019950 146491 bp DNA linear HTG 03-JAN-2000
DEFINITION Drosophila melanogaster. *** SEQUENCING IN PROGRESS ***.
ACCESSION AC019950
VERSION AC019950.1 GI:6664947
KEYWORDS HTG, HTGS PHASE2.
SOURCE Drosophila melanogaster (fruit fly)
ORGANISM Drosophila melanogaster (fruit fly)
Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
Ephydroidea; Drosophilidae; Drosophila.
1 (bases 1 to 146491)
REFERENCE Adams, M. and Venter, J.C.
AUTHORS

TITLE Direct Submission
JOURNAL Submitted (30-DEC-1999) Celera Genomics, 45 West Gude Drive,
Rockville, MD, USA
COMMENT This sequence was identified as CDM:10211502 by the submitter.
For more information on this record e-mail to fly@celera.com.
* NOTE: This is a 'working draft' sequence.
* This sequence will be replaced
* by the finished sequence as soon as it is available and
* the accession number will be preserved.

FEATURES
source
1. 146491
/organism="Drosophila melanogaster"
/mol_type="genomic DNA"
/db_xref="taxon:7227"

ORIGIN

Query Match 83.6%; Score 18.4; DB 2; Length 146491;
Best Local Similarity 95.0%; Pred. No. 7.2e+02;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 CGTCGTTTTGTCGCGTTT 21
38427 CGTCGTTTTGTCGCGTTT 38446

Db 38427 CGTCGTTTTGTCGCGTTT 38446

RESULT 15
LOCUS AC009904 182601 bp DNA linear INV 06-SEP-2001
DEFINITION Drosophila melanogaster, chromosome 3R, region 88B-88E, BAC clone
ACR32A03, complete sequence.
ACCESSION AC009904 AC007693
VERSION AC009904.7 GI:15451478
KEYWORDS HTG.
SOURCE Drosophila melanogaster (fruit fly)
ORGANISM Drosophila melanogaster
Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
Ephydroidea; Drosophilidae; Drosophila.

REFERENCE 1 (bases 1 to 182601)
AUTHORS Celinkier, S.E., Adams, M.D., Kronmiller, B., Tyler, D., Wan, K.H.,
Holt, R.A., Evans, C.A., Gocayne, J.D., Amanatides, P.G., Brannon, R.C.,
Rogers, Y., An, H., Baldwin, D., Bazon, J., Beeson, K.Y., Busam, D.A.,
Carlson, J.W., Center, A., Chape, M., Davenport, L.B., Dietz, S.M.,
Dodson, K., Dorsett, V., Doup, L.E., Doyle, C., Dresner, D., Farfan, D.,
Ferrera, S., Frise, E., Galle, R.F., Gary, N.S., George, R.A.,
Gonzalez, M., Houch, J., Hoskins, R.A., Hoshino, D., Howland, T.J.,
Ibegwam, C., Jalali, M., Kruse, D., Li, P., Mattei, B., Moshrefi, A.,
McIntosh, T.C., Moy, M., Murphy, B., Nelson, C., Nelson, K.A., Nunco, J.,
Paclet, J., Paragas, V., Park, S., Patel, S., Pfeiffer, B., Phouanavong, S.,
Pitman, G.S., Puri, V., Richards, S., Scheejer, F.,
Stapleton, M., Strong, R., Svirskas, R., Tector, C., Williams, S.M.,
Zaveri, J.S., Smith, H.O., Rubin, G.M. and Venter, J.C.

TITLE Sequencing of Drosophila chromosome 3R, region 88E-88E
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 182601)
AUTHORS Celinkier, S.E., Agbayani, A., Arcaina, T.T., Baxter, E., Blazej, R.G.,
Butenoff, C., Champe, M., Chavez, C., Chew, M., Ciesiolka, L.,
Doyle, C.M., Farfan, D.E., Galle, R., George, R.A., Harris, N.L.,
Hoskins, R.A., Houston, K.A., Hummasti, S.R., Karra, K., Kearney, L.,
Kim, E., Lee, B., Lewis, S., Li, P., Lomtan, M.A., Mazda, P.,
Moshrefi, A.R., Moshrefi, M., Nixon, K., Paclet, J.M., Park, S.,
Pfeiffer, B., Poon, L., Sequera, A., Sethi, H., Shit, E.,
Svirskas, R.R., Wan, K.H., Weinburg, T., Zhang, R., Zieran, L.L. and
Rubin, G.M.

TITLE Direct Submission
JOURNAL Submitted (06-SEP-1999) Drosophila Genome Center, Lawrence Berkeley
Laboratory, MS 64-121, Berkeley, CA 94720, USA
COMMENT On Sep 6, 2001 this sequence version replaced gi:13122705.
Sequence submitted by:
Berkeley Drosophila Genome Project
Lawrence Berkeley National Laboratory, MS 64-121
Berkeley, CA 94720
This sequence was assembled using end sequences from a whole genome

shotgun and from subclones of this BAC and its neighboring clones.
 For further information about this sequence, including its location
 and relationship to other sequences, please visit our sequence
 archive Web site (<http://www.fruitfly.org/sequence/>) or send email
 to bdg@fruitfly.berkeley.edu.

FEATURES

source

1. 182601

/organism="Drosophila melanogaster"

/mol_type="genomic DNA"

/strain="Y; cn bw sp"

/db_xref="taxon:7227"

/chromosome="3R"

/map="88E-88E"

/clone="BACR32A03 (D1083)"

/clone_id="RPCI-98 (Roswell Park Cancer Institute

Drosophila melanogaster BAC library, partial EcotI in
pBACe3.6)"

ORIGIN

Query Match 83.6%; Score 18.4; DB 3; Length 182601;

Best Local Similarity 95.0%; Pred. No. 7e+02;

Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 CGTCGTTTTCGTCGTTT 21

Db 115651 CGTCGTTTTCGTCGTTT 115670

Search completed: March 9, 2005, 18:45:34
 Job time : 3894 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using bw model

Run on: March 9, 2005, 17:17:45 ; Search time 3444 Seconds
(without alignments)
243.152 Million cell updates/sec

Title: US-10-613-228a-1

Perfect score: 22

Sequence: 1 tcgcgttttcgcgttttt 22

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 34239544 seqs, 19032134700 residues

Total number of hits satisfying chosen parameters: 6847908

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Database :

EST:*
1: gb_est1:*
2: gb_est2:*
3: gb_hsc:*
4: gb_est3:*
5: gb_est4:*
6: gb_est5:*
7: gb_est6:*
8: gb_gse1:*
9: gb_gse2:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
C 1	20.4	92.7	433	4	BG428594 602494726
C 2	19.4	88.2	957	9	AG126335 Pan tlog1
C 3	19.4	88.2	969	9	AG084147 Pan tlog1
C 4	19.4	86.4	473	7	CN197708 TGESTzy15
C 5	18.8	85.5	698	1	AV895065 AV895065
C 6	18.8	85.5	750	9	CL653482 PRI0118C
C 7	18.8	85.5	755	5	BP006348 BP006348
C 8	18.8	85.5	793	9	CL659802 PRI0135b
C 9	18.8	85.5	1173	3	CG745933 P039-1-A0
C 10	18.8	85.5	1201	3	CG704325 Tetradon
C 11	18.4	85.5	1372	9	CG751386 P045-4-D0
C 12	18.4	83.6	506	2	AV955361 AV955361
C 13	18.4	83.6	517	2	AV967757 AV967757
C 14	18.4	83.6	651	5	BM207212 BM207212
C 15	18.4	83.6	652	5	BM336085 BM336085
C 16	18.4	83.6	661	5	BM362385 BM362385
C 17	18.4	83.6	701	5	BP003979 BP003979
C 18	18.4	83.6	706	5	BM477298 BM477298
C 19	18.4	83.6	711	5	BM435032 BM435032
C 20	18.4	83.6	1101	9	CNS0001Y0 CNS0001Y0
C 21	18.4	81.8	236	7	W66251 W66251
C 22	17.8	80.9	246	6	CB365542 ZF001-P00
C 23	17.8	80.9	285	7	CN198693 TGESTzy16
C 24	17.8	80.9	342	6	CB353501 ZF001-P00

C 25	17.8	80.9	362	1	AV678562 AV678562
C 26	17.8	80.9	374	1	AV679629 AV679629
C 27	17.8	80.9	376	5	BM103452 BM103452
C 28	17.8	80.9	378	1	AV894777 AV894777
C 29	17.8	80.9	396	6	CB352916 CB352916
C 30	17.8	80.9	444	5	BM589623 BM589623
C 31	17.8	80.9	471	1	AV679587 AV679587
C 32	17.8	80.9	498	5	BM575333 BM575333
C 33	17.8	80.9	498	7	CN769410 CN769410
C 34	17.8	80.9	499	1	AV887894 AV887894
C 35	17.8	80.9	504	2	AM422790 AM422790
C 36	17.8	80.9	512	4	CG892389 CG892389
C 37	17.8	80.9	513	7	CN769717 CN769717
C 38	17.8	80.9	525	1	AV885063 AV885063
C 39	17.8	80.9	541	4	CG307575 CG307575
C 40	17.8	80.9	544	5	BM104367 BM104367
C 41	17.8	80.9	545	5	BM187581 BM187581
C 42	17.8	80.9	555	5	BQ420126 BQ420126
C 43	17.8	80.9	559	6	CB353813 CB353813
C 44	17.8	80.9	563	4	BT06183 BT06183
C 45	17.8	80.9	564	2	BE200845 BE200845

ALIGNMENTS

RESULT 1
BG428594/c 433 bp mRNA linear EST 14-MAR-2001
LOCUS 602494726P1 NIH_MGC_75 Homo sapiens CDNA clone IMAGE:4608342 5',
DEFINITION mRNA sequence.
ACCESSION BG428594 GI:13335100
VERSION BG428594.1 GI:13335100
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Cararrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 433)
NIH-MGC http://mgs.nci.nih.gov/.
AUTHORS National Institutes of Health, Mammalian Gene Collection (MGC)
TITLE Unpublished (1999)
JOURNAL
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cga@bbs-remail.nih.gov
Tissue Procurement: CLOUTCH Laboratories, Inc.
CDNA Library Preparation: CLOUTCH Laboratories, Inc.
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNU)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNU at:
http://image.llnl.gov
Plate: L1CM1350 row: d column: 07
High quality sequence stop: 132.
Location/Qualifiers
1. 433
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:4608342"
/lab host="DH10B (T1 phage-resistant)"
/clone.lib="NIH_MGC_75"
/note="Organ: Kidney; Vector: pDNR-LIB (Clontech); Site_1:
SfiI (ggccgctcgcc); Site_2: SfiI (ggccctatggcc); 5' and
3' adaptors were used in cloning as follows: 5' adaptor
sequence: 5'-CACGCGCATATGCGC-3' and 3' adaptor sequence:
5'-ATTCTAGAGCGGCGCGCGCATG-dt(30)BN-3' (where B = A,
C, or G and N = A, C, G, or T). Average insert size 1.65
kb (range 0.5-4.0 kb). 15/15 colonies contained inserts
by PCR. This library was enriched for full-length clones
and was constructed by Clontech Laboratories (Palo Alto,
CA). Note: this is a NIH_MGC Library."

ORIGIN

```

Query Match      92.7%; Score 20.4; DB 4; Length 433;
Best Local Similarity 95.5%; Pred. No. 1e+02;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY      1 TCGTCGTTTTTCGTCGCTTTT 22
      ||||| ||||| ||||| ||||| |||||
Db      390 TCGTCGTTTTTCGTCGCTTTT 369

RESULT 2
AGI26335/ 957 bp DNA linear GSS 04-NOV-2001
LOCUS      AGI26335
DEFINITION Pan troglodytes DNA, clone: PTB-136N20.F, genomic survey sequence.
ACCESSION  AGI26335
VERSION     AGI26335.1 GI:16655500
KEYWORDS    GSS.
SOURCE      Pan troglodytes (chimpanzee)
ORGANISM    Pan troglodytes
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Pan.

REFERENCE   1
AUTHORS     Fujiyama,A., Hattori,M., Toyoda,A., Taylor,T.D., Yada,T.,
            Totoki,Y., Watanabe,H. and Sakaki,Y.
TITLE       BAC end sequences of Library PTB
JOURNAL     Unpublished
AUTHORS      2 (bases 1 to 957)
            Fujiyama,A., Hattori,M., Toyoda,A., Taylor,T.D., Yada,T.,
            Totoki,Y., Watanabe,H. and Sakaki,Y.
TITLE       Direct Submission
SUBMITTED   (02-AUG-2001) Asao Fujiyama, The Institute of Physical
and Chemical Research (RIKEN), Genomic Sciences Center (GSC);
1-7-22 Suenhiro-chou,Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
(E-mail:chimps@sc.riken.go.jp, URL:http://hgp.gsc.riken.go.jp/,
Tel:81-45-503-9111, Fax:81-45-503-9170)
Clones are derived from the chimpanzee BAC library PTB This BAC end
was generated during the Rad process and may have higher chance of
clone tracking errors.
PRIMERS
Sequencing: -21M13

LIBRARY
Vector      : PKS145
R.Site 1    : SacI
R.Site 2    : SacI.
Location/Qualifiers
1..957
/organism="Pan troglodytes"
/mol_type="genomic DNA"
/db_xref="taxon:9598"
/clone="PTB-136N20.F"
/sex="male"
/cell_type="lymphoblast"
/clone_lib="PTB Chimpanzee Male BAC Library"

ORIGIN
Query Match      88.2%; Score 19.4; DB 9; Length 957;
Best Local Similarity 95.2%; Pred. No. 2.8e+02;
Matches 20; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY      1 TCGTCGTTTTTCGTCGCTTTT 21
      ||||| ||||| ||||| ||||| |||||
Db      795 TCGTCGTTTTTCGTCGCTTTT 775

RESULT 3
AG084147 969 bp DNA linear GSS 03-NOV-2001
LOCUS      AG084147
DEFINITION Pan troglodytes DNA, clone: PTB-081M22.F, genomic survey sequence.
ACCESSION  AG084147
VERSION     AG084147.1 GI:16635949
KEYWORDS    GSS.
SOURCE      Pan troglodytes (chimpanzee)
ORGANISM    Pan troglodytes
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

```

```

REFERENCE   1
AUTHORS     Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Pan.
            Fujiyama,A., Hattori,M., Toyoda,A., Taylor,T.D., Yada,T.,
            Totoki,Y., Watanabe,H. and Sakaki,Y.
TITLE       BAC end sequences of Library PTB
JOURNAL     Unpublished
AUTHORS      2 (bases 1 to 969)
            Fujiyama,A., Hattori,M., Toyoda,A., Taylor,T.D., Yada,T.,
            Totoki,Y., Watanabe,H. and Sakaki,Y.
TITLE       Direct Submission
SUBMITTED   (02-AUG-2001) Asao Fujiyama, The Institute of Physical
and Chemical Research (RIKEN), Genomic Sciences Center (GSC);
1-7-22 Suenhiro-chou,Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
(E-mail:chimps@sc.riken.go.jp, URL:http://hgp.gsc.riken.go.jp/,
Tel:81-45-503-9111, Fax:81-45-503-9170)
Clones are derived from the chimpanzee BAC library PTB This BAC end
was generated during the Rad process and may have higher chance of
clone tracking errors.
PRIMERS
Sequencing: -21M13

LIBRARY
Vector      : PKS145
R.Site 1    : SacI
R.Site 2    : SacI.
Location/Qualifiers
1..969
/organism="Pan troglodytes"
/mol_type="genomic DNA"
/db_xref="taxon:9598"
/clone="PTB-081M22.F"
/sex="male"
/cell_type="lymphoblast"
/clone_lib="PTB Chimpanzee Male BAC Library"

ORIGIN
Query Match      88.2%; Score 19.4; DB 9; Length 969;
Best Local Similarity 95.2%; Pred. No. 2.8e+02;
Matches 20; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY      2 CGTCGTTTTTCGTCGCTTTT 22
      ||||| ||||| ||||| ||||| |||||
Db      714 CGTCGTTTTTCGTCGCTTTT 734

RESULT 4
CN197708 473 bp mRNA linear EST 05-APR-2004
LOCUS      CN197708
DEFINITION TgESTzy155a11.y1 TgVEG118 Tachyzoite CDNA Library-2 Toxoplasma
gondii cDNA clone TgESTzy155a11.y1 5', mRNA sequence.
ACCESSION  CN197708
VERSION     CN197708.1 GI:46222647
KEYWORDS    EST.
SOURCE      Toxoplasma gondii
            Toxoplasma gondii
            Eukaryota; Alveolata; Apicomplexa; Coccidia; Eimeriida;
            Sarcocystidae; Toxoplasma.
ORGANISM    Sarcocystidae; Toxoplasma.

REFERENCE   1 (bases 1 to 473)
AUTHORS     Tang,K., Cole,R., Fogarty,S., Sibley,L.D., Ajioke,J.A., White,M.,
            Clifton,S., Pape,D., Martin,J., Wylie,T., Dante,M., Marra,M.,
            Hillier,L., Kucaba,T., Theising,B., Bowers,Y., Gibbons,M.,
            Ritzer,E., Bennett,J., Franklin,C., Tsagarisvilli,R., Ronko,I.,
            Kennedy,S., Maguire,L., Waterston,R. and Wilson,R.
TITLE       Toxoplasma EST Project
JOURNAL     Unpublished (2001)
COMMENT     Contact: Clifton, S.
            Toxoplasma EST Project
            Washington University School of Medicine
            4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA
            Tel: 314 286 1800
            Fax: 314 286 1810
            Email: toxo@watson.wustl.edu
            Contact David Sibley (toxost@borcim.wustl.edu) for further
            information relating to organism, libraries, or clone availability.

```

Seq primer: -40UP from Glibco
High quality sequence stop: 473.
Location/Qualifiers
1..473

/organism="Toxoplasma gondii"
/mol_type="mRNA"
/db_xref="taxon:5811"
/clone="TGSTz155a1.y1"
/dev_stage="Tachyzoite"
/lab_host="GC10"
/clone_lib="TVEG118 Tachyzoite cDNA library-2"
/note="Vector: pBluescript SK, Site 1: EcoRI, Site 2: XhoI; The library was constructed by Kelang Tang, Robert Cole and L. David Sibley at Washington University. cDNAs were synthesized from poly(A)+ RNasy oligo d(T) priming, size-selected and directionally cloned into the Uni-ZAP XR lambda vector (Stratagene). The primary library was mass excised as phagemids and rescued in SOLR cells. The plasmid library was recovered from the SOLR cells and transformed in mass into GC10 cells for sequencing.
WARNING: the library may contain a small percentage contaminants from human fibroblast cells."

ORIGIN

Query Match 86.4%; Score 19; DB 7; Length 473;
Best Local Similarity 100.0%; Pred. No. 4.2e+02;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Cy 3 GTCGTTTTCGTGGCTTT 21
|||||
Db 153 CTCGTTTTCGTGGCTTT 171

RESULT 5
AV895065/c 698 bp mRNA linear EST 09-NOV-2001
LOCUS
DEFINITION AV895065 Nori Satoh unpublished cDNA library, young adult Clona
intestinalis cDNA clone rciad40014 3', mRNA sequence.
ACCESSION
AV895065
VERSION
AV895065.1 GI:16884161
KEYWORDS
EST.
SOURCE
Clona intestinalis
ORGANISM
Clona intestinalis
Eukaryota; Metazoa; Chordata; Urochordata; Ascidiacea; Enterogona;
Phlebobranchia; Cloniidae; Clona.
1 (bases 1 to 698)
Satch,N., Satou,Y., Kohara,Y. and Shin-I,T.
Expressed genes in Clona intestinalis
JOURNAL
Unpublished (2000)
COMMENT
Contact: Nori Satoh
Department of Zoology
Sakyo-ku, Kyoto, Kyoto 606-8502, Japan
Tel: 81-75-753-4081
Fax: 81-75-705-1113
Email: sato@ascidian.zool.kyoto-u.ac.jp.
Location/Qualifiers

FEATURES
SOURCE
1..698
/organism="Clona intestinalis"
/mol_type="mRNA"
/db_xref="taxon:7719"
/clone="rciad40014"
/tissue_type="whole animal"
/dev_stage="young adult"
/clone_lib="Nori Satoh unpublished cDNA library, young adult"

ORIGIN

Query Match 85.5%; Score 18.8; DB 1; Length 698;
Best Local Similarity 90.9%; Pred. No. 5.2e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Cy 1 TCGTGTTTTCGTGGCTTT 22
|||||
Db 484 TCGTGTTTTCGTGGCTTT 463

RESULT 6
CL653482 750 bp DNA linear GSS 09-JUL-2004
LOCUS
DEFINITION PR10118c.G01 - PR10118c.B21 (750) Mixed stage fosmid library of P.
pacificus var. California Pristionchus pacificus genomic, genomic
survey sequence.
ACCESSION
CL653482
VERSION
CL653482.1 GI:50132352
KEYWORDS
GSS.
SOURCE
Pristionchus pacificus
ORGANISM
Pristionchus pacificus
Eukaryota; Metazoa; Nematoda; Chromadorea; Diplogasterida;
Neodiplogasteridae; Pristionchus.
1 (bases 1 to 750)
Srinivasan,J., Otto,G.W., Kahlow,U., Geisler,R. and Sommer,R.J.
Appar: an Acedb database for the nematode satellite organism
Pristionchus pacificus
Nucleic Acids Res. 32 (1), D421-D422 (2004)
CONTACT: Sommer RJ
Evolutionary Biology
Max-Planck-Institute for Developmental Biology
Spemannstr. 37-39, Tuebingen D-72076, Germany
Tel: 00497071601371
Fax: 00497071601498
Email: ralf.sommer@uebingen.mpg.de
This library was generated at Caltech, Pasadena, USA and end
sequenced at Vancouver, Canada.
Seq primer: T7
Class: fosmid ends.
Location/Qualifiers

FEATURES
SOURCE
1..750
/organism="Pristionchus pacificus"
/mol_type="genomic DNA"
/strain="California"
/db_xref="taxon:54126"
/clone_lib="Mixed stage fosmid library of P. pacificus
var. California"
/note="Vector: pBpifos-5 Fosmid vector"

ORIGIN

Query Match 85.5%; Score 18.8; DB 9; Length 750;
Best Local Similarity 90.9%; Pred. No. 5.1e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Cy 1 TCGTGTTTTCGTGGCTTT 22
|||||
Db 457 TCGTGTTTTCGTGGCTTT 478

RESULT 7
BP006348 755 bp mRNA linear EST 15-MAR-2002
LOCUS
DEFINITION BP006348 Nori Satoh unpublished cDNA library, young adult Clona
intestinalis cDNA clone ciad40014 5', mRNA sequence.
ACCESSION
BP006348
VERSION
BP006348.1 GI:19497825
KEYWORDS
EST.
SOURCE
Clona intestinalis
ORGANISM
Clona intestinalis
Eukaryota; Metazoa; Chordata; Urochordata; Ascidiacea; Enterogona;
Phlebobranchia; Cloniidae; Clona.
1 (bases 1 to 755)
Satch,N., Satou,Y., Kohara,Y. and Shin-I,T.
Expressed genes in Clona intestinalis
JOURNAL
Unpublished (2000)
COMMENT
Contact: Nori Satoh
Department of Zoology
Kyoto University

REFERENCE
Satch,N., Satou,Y., Kohara,Y. and Shin-I,T.
Expressed genes in Clona intestinalis
JOURNAL
Unpublished (2000)
CONTACT: Nori Satoh
Department of Zoology
Kyoto University

Sakyo-ku, Kyoto, Kyoto 606-8502, Japan
Tel: 81-75-753-4081
Fax: 81-75-705-1113
Email: satcho@sci.dian.zool.kyoto-u.ac.jp.

FEATURES

source

Location/Qualifiers
1..755

/organism="Clona intestinalis"
/mol_type="mRNA"
/db_xref="taxon:7719"
/clone="clad40014"
/issue_type="whole animal"
/dev_stage="young adult"
/clone_lib="Nori Satoh unpublished cDNA library, young adult"

ORIGIN

Query Match 85.5%; Score 18.8; DB 5; Length 755;
Best Local Similarity 90.9%; Pred. No. 5.1e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY

1 TCGTCGTTTTCGTGCGTTT 22
|||||
277 TCGTCGTTTTCGTGCGTTT 298

RESULT 8 793 bp DNA linear GSS 09-JUL-2004
CL659802 PRI0135D_D02 - PRI0135B.B21 (793) Mixed stage fosmid library of P.
LOCUS pacificus var. California Pristionchus pacificus genomic, genomic
DEFINITION survey sequence.

ACCESSION CL659802 GI:50144272
VERSION CL659802
KEYWORDS GSS.

SOURCE

Pristionchus pacificus
Pristionchus pacificus
Eukaryota; Metazoa; Nematoda; Chromadorea; Diplogasterida;
Neodiplogasteridae; Pristionchus.

ORGANISM

1 (bases 1 to 793)
Srinivasan,J., Otto,G.W., Kahlow,U., Geisler,R. and Sommer,R.J.
AppADB: an AceDB database for the nematode satellite organism
Pristionchus pacificus
Nucleic Acids Res. 32 (1), D421-D422 (2004)

REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT

Max-Planck-Institute for Developmental Biology
Spemannstr. 37-39, Tuebingen D-72076, Germany
Tel: 00497071601371
Fax: 00497071601498
Email: ralf.sommer@tuebingen.mpg.de
This library was generated at Caltech, Pasadena, USA and end
sequenced at Vancouver, Canada.
Seq primer: T7
Class: fosmid ends.

FEATURES
source
Location/Qualifiers
1..793

/organism="Pristionchus pacificus"
/mol_type="genomic DNA"
/strain="California"
/db_xref="taxon:54126"
/clone_lib="Mixed stage fosmid library of P. pacificus
var. California"
/note="Vector: pEplfos-5 Fosmid vector"

ORIGIN

Query Match 85.5%; Score 18.8; DB 9; Length 793;
Best Local Similarity 90.9%; Pred. No. 5.1e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY

1 TCGTCGTTTTCGTGCGTTT 22
|||||
466 TCGTCGTTTTCGTGCGTTT 487

RESULT 9 1173 bp DNA linear GSS 24-OCT-2003
CG745933/c P039-1-A09.za Ppa EcORI BAC library Pristionchus pacificus genomic,
LOCUS genomic survey sequence.

DEFINITION

ACCESSION CG745933 GI:37966859
VERSION CG745933
KEYWORDS GSS.

SOURCE

Pristionchus pacificus
Pristionchus pacificus
Eukaryota; Metazoa; Nematoda; Chromadorea; Diplogasterida;
Neodiplogasteridae; Pristionchus.

ORGANISM

1 (bases 1 to 1173)
Srinivasan,J., Sinz,W., Jesse,T., Wiggers-Perebolte,L., Jansen,K.,
Bunjer,J., van der Meulen,M. and Sommer,R.J.
An integrated physical and genetic map of the nematode Pristionchus
pacificus
Mol. Genet. Genomics 269 (5), 715-722 (2003)

REFERENCE

AUTHORS
TITLE
JOURNAL
MEDLINE
PUBMED
COMMENT

Evolutionary Biology
Max-Planck-Institute for Developmental Biology
Spemannstr. 37-39, Tuebingen D-72076, Germany
Tel: 00497071601371
Fax: 00497071601498
Email: ralf.sommer@tuebingen.mpg.de
Class: BAC ends.

FEATURES
source
Location/Qualifiers
1..1173

/organism="Pristionchus pacificus"
/mol_type="genomic DNA"
/strain="California"
/db_xref="taxon:54126"
/clone_lib="Ppa EcORI BAC library"
/note="The library was generated by a partial digest of
the genomic DNA with EcORI and cloning into the BAC
vector."

ORIGIN

Query Match 85.5%; Score 18.8; DB 9; Length 1173;
Best Local Similarity 90.9%; Pred. No. 5.1e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY

1 TCGTCGTTTTCGTGCGTTT 22
|||||
164 TCGTCGTTTTCGTGCGTTT 143

RESULT 10 1201 bp mRNA linear HTC 19-AUG-2004
CR704325 Tetraodon nigroviridis full-length cDNA.
LOCUS CR704325
DEFINITION CR704325.1 GI:51202234
ACCESSION CR704325
VERSION CR704325.1
KEYWORDS HTC; cDNA; full-length; Tetraodon nigroviridis.

SOURCE

Tetraodon nigroviridis
Tetraodon nigroviridis
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
Acanthomorpha; Acanthopterygii; Percomorpha; Tetraodontiformes;
Tetraodontidae; Tetraodontidae; Tetraodon.

ORGANISM

1 (bases 1 to 1201)
Genoscope.
Direct Submission
Submitted (10-AUG-2004) Genoscope - Centre National de Sequencage -
2 rue Gaston Cremieux, CP 5706 - 91057 Evry cedex - FRANCE
(E-mail : sequef@genoscope.cns.fr - Web : www.genoscope.cns.fr)
The sequences are based on single pass reads.
More information available at
http://www.genoscope.cns.fr/tetraodon.

REFERENCE

AUTHORS
TITLE
JOURNAL
COMMENT

FEATURES
Location/Qualifiers

source
1. .1201
/organism="Tetracodon nigroviridis"
/mol_type="mRNA"
/db_xref="taxon:99883"
/tissue_type="Eyes"

ORIGIN
Query Match 85.5%; Score 18.8; DB 3; Length 1201;
Best Local Similarity 90.9%; Pred. No. 5.1e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 TCGTCGTTTTCGTCGTTT 22
|||||
Db 558 TCGTCGTTTTCGTCGTTT 579

RESULT 11
CG751386 1372 bp DNA linear GSS 24-OCT-2003
LOCUS P045-4-D09.yb Ppa EcORI BAC Library Pristionchus pacificus genomic.
DEFINITION genomic survey sequence.
ACCESSION CG751386
VERSION CG751386.1 GI:37973790
KEYWORDS GSS.
SOURCE Pristionchus pacificus
ORGANISM Pristionchus pacificus
Eukaryota; Metazoa; Nematoda; Chromadorea; Diplogasterida;
Neodiplogasteridae; Pristionchus.
REFERENCE 1 (bases 1 to 1372)
AUTHORS Striavaan,U., Sinz,W., Jesse,T., Wigers-Perebolte,L., Jansen,K.,
Buntjer,J., van der Meulen,M. and Sommer,R.J.
TITLE An integrated physical and genetic map of the nematode Pristionchus
JOURNAL Mol. Genet. Genomics 269 (5), 715-722 (2003)
MEDLINE 22835951
PUBMED 12884007
COMMENT Contact: Sommer RJ
Evolutionary Biology
Max-Planck-Institute for Developmental Biology
Spemannstr. 37-39, Tuebingen D-72076, Germany
Tel: 00497071601371
Fax: 00497071601498
Email: ralf.sommer@tuebingen.mpg.de
Class: BAC ends
Location/Qualifiers
1. .1372
/organism="Pristionchus pacificus"
/mol_type="genomic DNA"
/strain="California"
/db_xref="taxon:54126"
/clone_id="Ppa EcORI BAC library"
/note="The library was generated by a partial digest of
the genomic DNA with EcORI and cloning into the BAC
vector."

ORIGIN
Query Match 85.5%; Score 18.8; DB 9; Length 1372;
Best Local Similarity 90.9%; Pred. No. 5.1e+02;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 TCGTCGTTTTCGTCGTTT 22
|||||
Db 468 TCGTCGTTTTCGTCGTTT 447

RESULT 12
AV955361 506 bp mRNA linear EST 14-MAR-2002
LOCUS AV955361 Nori Satoh unpublished cDNA library, egg Ciona
DEFINITION intestinalis cDNA clone cleg08a22 5', mRNA sequence.
ACCESSION AV955361
VERSION AV955361.1 GI:19443660
KEYWORDS EST.

SOURCE
Ciona intestinalis
ORGANISM Ciona intestinalis
Eukaryota; Metazoa; Chordata; Urochordata; Ascidiacea; Enterogona;
Phlebobranchia; Cionidae; Ciona.
REFERENCE 1 (bases 1 to 506)
AUTHORS Satoh,N., Satou,Y., Kohara,Y. and Shin-i,T.
TITLE Expressed genes in Ciona intestinalis
JOURNAL Unpublished (2000)
COMMENT Contact: Nori Satoh
Department of Zoology
Kyoto University
Sakyo-ku, Kyoto, Kyoto 606-8502, Japan
Tel: 81-75-753-4081
Fax: 81-75-705-1113
Email: satoh@ascidian.zool.kyoto-u.ac.jp.
Location/Qualifiers
1. .506
/organism="Ciona intestinalis"
/mol_type="mRNA"
/db_xref="taxon:7719"
/clone="cleg08a22"
/tissue_type="whole animal"
/dev stage="egg"
/clone_id="Nori Satoh unpublished cDNA library, egg"

ORIGIN
Query Match 83.6%; Score 18.4; DB 2; Length 506;
Best Local Similarity 95.0%; Pred. No. 7.8e+02;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TCGTCGTTTTCGTCGTTT 20
|||||
Db 125 TCGTCGTTTTCGTCGTTT 106

RESULT 13
AV967757 517 bp mRNA linear EST 14-MAR-2002
LOCUS AV967757 Nori Satoh unpublished cDNA library, egg Ciona
DEFINITION intestinalis cDNA clone cleg23e07 5', mRNA sequence.
ACCESSION AV967757
VERSION AV967757.1 GI:19457453
KEYWORDS EST.
SOURCE Ciona intestinalis
ORGANISM Ciona intestinalis
Eukaryota; Metazoa; Chordata; Urochordata; Ascidiacea; Enterogona;
Phlebobranchia; Cionidae; Ciona.
REFERENCE 1 (bases 1 to 517)
AUTHORS Satoh,N., Satou,Y., Kohara,Y. and Shin-i,T.
TITLE Expressed genes in Ciona intestinalis
JOURNAL Unpublished (2000)
COMMENT Contact: Nori Satoh
Department of Zoology
Kyoto University
Sakyo-ku, Kyoto, Kyoto 606-8502, Japan
Tel: 81-75-753-4081
Fax: 81-75-705-1113
Email: satoh@ascidian.zool.kyoto-u.ac.jp.
Location/Qualifiers
1. .517
/organism="Ciona intestinalis"
/mol_type="mRNA"
/db_xref="taxon:7719"
/clone="cleg23e07"
/tissue_type="whole animal"
/dev stage="egg"
/clone_id="Nori Satoh unpublished cDNA library, egg"

ORIGIN
Query Match 83.6%; Score 18.4; DB 2; Length 517;
Best Local Similarity 95.0%; Pred. No. 7.8e+02;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TCGTCGTTTTCGTGCGTTT 20
| | | | | | | | | | | | | | | | | | | | | | | | | |
Db 117 TCGTCGTCCTTTCGTGCGTTT 98

RESULT 14

BW207212/c 651 bp mRNA linear EST 05-NOV-2002
LOCUS BW207212 Nori Satoh unpublished cDNA library, cleaving embryo Ciona
DEFINITION intestinalis cDNA clone cici097a05 5', mRNA sequence.
ACCESSION BW207212
KEYWORDS EST.
SOURCE GI:24621826
ORGANISM Ciona intestinalis

REFERENCE

AUTHORS 1 TCGTCGTTTTCGTGCGTTT 20
TITLE | | | | | | | | | | | | | | | | | | | | | | | | | |
JOURNAL / | | | | | | | | | | | | | | | | | | | | | | | | | |
COMMENT / | | | | | | | | | | | | | | | | | | | | | | | | | |
Department of Zoology
Kyoto University
Sakyo-ku, Kyoto, Kyoto 606-8502, Japan
Tel: 81-75-753-4081
Fax: 81-75-705-1113
Email: satcho@ascidian.zool.kyoto-u.ac.jp.
Location/Qualifiers

FEATURES

1.651
/organism="Ciona intestinalis"
/mol_type="mRNA"
/db_xref="taxon:7719"
/clone="cici097a05"
/tissue_type="whole body"
/dev_stage="cleaving embryo"
/clone_lib="Nori Satoh unpublished cDNA library, cleaving embryo"

ORIGIN

Query Match 83.6%; Score 18.4; DB 5; Length 651;
Best Local Similarity 95.0%; Pred.No.7.8e+02;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TCGTCGTTTTCGTGCGTTT 20
| | | | | | | | | | | | | | | | | | | | | | | | | |
Db 612 TCGTCGTCCTTTCGTGCGTTT 593

RESULT 15
BW336085/c 652 bp mRNA linear EST 27-MAY-2004
LOCUS BW336085 Yutaka Satou unpublished cDNA library, embryo whole animal
DEFINITION Ciona intestinalis cDNA clone ciem801019 5', mRNA sequence.
ACCESSION BW336085
KEYWORDS EST.
SOURCE GI:47747886
ORGANISM Ciona intestinalis

Eukaryota; Metazoa; Chordata; Urochordata; Ascidiacea; Enterogona;
Phlebobranchia; Clonidae; Ciona.
1 (bases 1 to 652)
Satou,Y., Shin-I., Kohara,Y. and Satoh,N.
Expressed genes in Ciona intestinalis (2004)
Unpublished (2004)
Contact: Yutaka Satou
Department of Zoology
Kyoto University
Sakyo-ku, Kyoto, Kyoto 606-8502, Japan
Tel: 81-75-753-4095
Fax: 81-75-705-1113
Email: yutaka@ascidian.zool.kyoto-u.ac.jp.
Location/Qualifiers

REFERENCE

AUTHORS 1 TCGTCGTTTTCGTGCGTTT 20
TITLE | | | | | | | | | | | | | | | | | | | | | | | | | |
JOURNAL / | | | | | | | | | | | | | | | | | | | | | | | | | |
COMMENT / | | | | | | | | | | | | | | | | | | | | | | | | | |
Department of Zoology
Kyoto University
Sakyo-ku, Kyoto, Kyoto 606-8502, Japan
Tel: 81-75-753-4095
Fax: 81-75-705-1113
Email: yutaka@ascidian.zool.kyoto-u.ac.jp.
Location/Qualifiers

FEATURES

source 1.652
/organism="Ciona intestinalis"
/mol_type="mRNA"
/db_xref="taxon:7719"
/clone="ciem801019"
/tissue_type="whole animal"
/dev_stage="embryo"
/clone_lib="Yutaka Satou unpublished cDNA library, embryo whole animal"

ORIGIN

Query Match 83.6%; Score 18.4; DB 5; Length 652;
Best Local Similarity 95.0%; Pred.No.7.8e+02;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TCGTCGTTTTCGTGCGTTT 20
| | | | | | | | | | | | | | | | | | | | | | | | | |
Db 406 TCGTCGTCCTTTCGTGCGTTT 387

Search completed: March 9, 2005, 19:45:52
Job time : 3450 secs

CC fungicide, and antiparasitic activity. A composition may act as an
CC interleukin antagonist-4, or interleukin antagonist-5, and may have a use
CC in gene therapy. The methods and compositions of the present invention
CC are useful for diagnosing, preventing and/or treating infectious disease,
CC allergy, asthma, cancer, where the infectious disease is a herpes simplex
CC virus, bacterial, fungal or parasitic infection, and where the cancer is
CC a biliary tract cancer, bone cancer, brain and CNS cancer, breast cancer,
CC cervical cancer, choriocarcinoma, colon cancer, connective tissue cancer,
CC endometrial cancer, esophageal cancer, eye cancer, gastric cancer,
CC Hodgkin's lymphoma, intraepithelial neoplasms, larynx cancer, lymphomas,
CC liver cancer, lung cancer (e.g. small cell and non-small cell), melanoma,
CC neuroblastoma, oral cavity cancer, ovarian cancer, pancreas cancer,
CC prostate cancer, rectal cancer, sarcomas, skin cancer, testicular cancer,
CC thyroid cancer and renal cancer. The present sequence represents an
CC immunostimulatory nucleic acid molecule of the invention.
XX
SQ Sequence 22 BP; 0 A; 4 C; 5 G; 13 T; 0 U; 0 Other;

Query Match 100.0%; Score 22; DB 12; Length 22;
Best Local Similarity 100.0%; Pred. No. 5.3;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCGTCGTTTTTCGTGCGTTTTT 22
Db 1 TCGTCGTTTTTCGTGCGTTTTT 22

RESULT 2
ADK19243
ID ADK19243 standard; DNA; 22 BP.
XX
AC ADK19243;
XX
DT 20-MAY-2004 (first entry)
XX
DE Immunostimulatory nucleic acid #289.
XX
KW Immunostimulatory nucleic acid; asthma; allergy; cancer;
KW infectious disease; autoimmune disease; airway remodeling;
KW chronic obstructive pulmonary disease; asthma; IL-6; interleukin-6;
KW TNFalpha; tumour necrosis factor alpha; IFNalpha; interferon-alpha;
KW IFNgamma; interferon-gamma; IP-10; interferon inducible protein;
KW viral infection; bacteria infection; parasitic infection; ss.
XX
OS Synthetic.
XX
PN WO2004016805-A2.
XX
PD 26-FEB-2004.
XX
PF 19-AUG-2003; 2003WO-US025935.
XX
PR 19-AUG-2002; 2002US-0404479P.
PR 19-AUG-2002; 2002US-0404820P.
PR 27-NOV-2002; 2002US-0429701P.
PR 14-FEB-2003; 2003US-0447377P.
XX
XX (COLE-) COLEY PHARM GROUP INC.
PA (COLE-) COLEY PHARM GMBH.
XX
PI Krieg AM, Samulowitz U, Vollmer J, Uhlmann E, Jurk M, Lipford G,
PI Rankin R;
XX
DR WPI; 2004-257200/24.
XX
XX New immunostimulatory nucleic acid molecule having pyrimidine-purine
PT dinucleotide and a chimeric backbone, useful in treating and preventing
PT asthma, allergy, cancer, infectious disease, autoimmune disease or airway
PT remodeling.
XX
PS Example 15; SEQ ID NO 290; 276pp; English.
XX
CC The invention relates to an immunostimulatory nucleic acid molecule

CC comprising an internal pyrimidine-purine (YZ) dinucleotide and chimeric
CC backbone, where one internal YZ dinucleotide has a phosphodiester(-like)
CC internucleotide linkage, where optionally each additional internal YZ
CC dinucleotide has a phosphodiester(-like) or stabilised internucleotide
CC linkage, where other internucleotide linkages are stabilised. The
CC oligonucleotide is useful in stimulating or modulating an immune
CC response. The medicament shifts the immune response to a Th1 biased
CC response from a Th2 biased response. The oligonucleotide is also useful
CC in the manufacture of a medicament for treating asthma, allergy, cancer,
CC infectious disease, autoimmune disease, airway remodeling or chronic
CC obstructive pulmonary disease or in treating a subject who is a smoker or
CC who is free of symptoms of asthma. The oligonucleotide is useful in
CC inducing cytokine expression, e.g. IL-6 (interleukin-6), TNFalpha (tumour
CC necrosis factor alpha), IFNalpha (interferon-alpha), IFNgamma (interferon
CC -gamma) and IP-10 (interferon inducible protein). The oligonucleotide is
CC also useful in treating and preventing infections caused by viruses,
CC bacteria and parasites. The present sequence represents an
CC immunostimulatory nucleic acid.
XX
SQ Sequence 22 BP; 0 A; 4 C; 5 G; 13 T; 0 U; 0 Other;

Query Match 100.0%; Score 22; DB 12; Length 22;
Best Local Similarity 100.0%; Pred. No. 5.3;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCGTCGTTTTTCGTGCGTTTTT 22
Db 1 TCGTCGTTTTTCGTGCGTTTTT 22

RESULT 3
ADO4307
ID ADO4307 standard; DNA; 22 BP.
XX
AC ADO4307;
XX
DT 29-JUL-2004 (first entry)
XX
DE Nucleotide sequence of a CpG ODN of class B.
XX
KW HCV infection; CpG therapy; immunostimulatory; hepatotropic; virucide;
KW gene therapy; ss.
XX
OS Synthetic.
XX
PN WO2004039829-A2.
XX
PD 13-MAY-2004.
XX
PF 29-OCT-2003; 2003WO-IB005520.
XX
PR 29-OCT-2002; 2002US-0421987P.
XX
PA (COLE-) COLEY PHARM GROUP LTD.
PA (COLE-) COLEY PHARM GMBH.
XX
XX Ahluwalia NK, Eflier SM, Davis HU, Vollmer J;
PI
XX
DR WPI; 2004-376156/35.
XX
PT Treating a patient having hepatitis C virus (HCV) infection that was not
PT successfully treated using a previous non-CpG therapy comprises
PT administering to a subject a CpG immunostimulatory nucleic acid.
XX
PS Example; SEQ ID NO 6; 89pp; English.
XX
XX The invention relates to creating a patient having hepatitis C virus
CC (HCV) infection that was not successfully treated using a previous non-
CC CpG therapy. The method involves administering to a subject in need of
CC such treatment a CpG immunostimulatory nucleic acid in an amount
CC effective to treat the infection. In the treatment method, the non-CpG
CC therapy includes interferon-alpha. The interferon-alpha is interferon-
CC alpha-2b, interferon-alpha-2a or consensus interferon-alpha. The non-CpG

CC therapy includes interferon-alpha or pegylated interferon-alpha and
CC ribavirin. The CpG immunostimulatory nucleic acid is A, B or C class CpG
CC immunostimulatory nucleic acid. The method further comprises
CC administering interferon-alpha to the subject. The interferon-alpha is
CC administered substantially simultaneously with the CpG immunostimulatory
CC nucleic acid. The CpG immunostimulatory nucleic acid comprises a backbone
CC modification, preferably a phosphorothionate backbone modification. The
CC CpG immunostimulatory nucleic acid comprises a semi-soft backbone. The
CC method is useful for treating a patient having hepatitis C virus (HCV)
CC infection that was not successfully treated using a previous non-CpG
CC therapy. Sequences ADO44302-ADO44317 represent examples of CpG
CC oligodeoxynucleotides (ODN) which were used in the experiments to
CC exemplify the methods of the invention.

XX Sequence 22 BP; 0 A; 4 C; 5 G; 13 T; 0 U; 0 Other;
SQ

Query Match 100.0%; Score 22; DB 12; Length 22;
Best Local Similarity 100.0%; Pred. No. 5.3;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCGTCGTTTTCGTCGTTT 22
1 TCGTCGTTTTCGTCGTTT 22
DB

RESULT 4
ABQ47667/c
ID ABQ47667 standard; DNA; 618 BP.
XX
AC ABQ47667;
XX
DT 12-JUL-2002 (first entry)
XX
DE Oligonucleotide for detecting cytosine methylation SEQ ID NO 34258.
XX
KW Human; cytosine methylation; 5'-CpG-3'; uracil; cytosine; diagnosis;
KW drug; side effect; cancer; central nervous system; cardiovascular;
KW gastrointestinal; respiratory system; single nucleotide polymorphism;
KW SNP; cell differentiation; ds.
XX
OS Homo sapiens.
XX
PN WO200218632-A2.
XX
PD 07-MAR-2002.
XX
PF 01-SEP-2001; 2001WO-EP010074.
XX
PR 01-SEP-2000; 2000DE-01043826.
PR 05-SEP-2000; 2000DE-01044543.
XX
PA (EPIC-) EPIGENOMICS AG.
XX
PI Olek A, Piepenbrock C, Berlin K, Guetig D;
XX
DR WPI; 2002-371829/40.
XX
PT Determining the degree of cytosine methylation in genomic DNA, useful for
PT diagnosis and prognosis, comprises selective hybridization of amplicons
PT from chemically treated DNA.
XX
PS Claim 12; 56pp + Sequence Listing; 56pp; German.

CC This invention describes a novel method for determining the degree of
CC methylation of a particular cytosine in a motif 5'-CpG-3', present in a
CC genomic sample of DNA. The sample is treated chemically to convert
CC cytosine (C) but not methylated C, to uracil, then part of the genomic
CC DNA that contains the target C is amplified to form a labeled amplicon.
CC The amplicon is hybridised to two classes, each with at least one member,
CC of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers and the
CC degree of hybridisation to both classes is determined from the label on
CC the amplicon. From the ratio of labels hybridised to the two classes of
CC oligomers, the degree of methylation is calculated. The method is used:

X

CC (i) for diagnosis and/or prognosis of side effects of therapeutic drugs
CC and of a wide range of diseases, e.g. cancer, disorders of the central
CC nervous, cardiovascular, gastrointestinal and respiratory systems etc.,
CC particularly by detecting mutations or single nucleotide polymorphisms
CC (SNP's) and (ii) for differentiation of cell or tissue types and for
CC investigating cell differentiation. The method allows the methylation
CC status of many C residues to be determined simultaneously. ABQ1410
CC ABQ54121 represent genomic DNA sequences used to illustrate the method
CC for determining the degree of cytosine methylation described in the
CC disclosure of the invention

XX Sequence 618 BP; 241 A; 204 C; 58 G; 115 T; 0 U; 0 Other;
SQ

Query Match 92.7%; Score 20.4; DB 6; Length 618;
Best Local Similarity 95.5%; Pred. No. 26;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TCGTCGTTTTCGTCGTTT 22
521 TCGTCGTTTTCGTCGTTT 500
DB

RESULT 5
ABQ47666
ID ABQ47666 standard; DNA; 618 BP.
XX
AC ABQ47666;
XX
DT 12-JUL-2002 (first entry)
XX
DE Oligonucleotide for detecting cytosine methylation SEQ ID NO 34257.
XX
KW Human; cytosine methylation; 5'-CpG-3'; uracil; cytosine; diagnosis;
KW drug; side effect; cancer; central nervous system; cardiovascular;
KW gastrointestinal; respiratory system; single nucleotide polymorphism;
KW SNP; cell differentiation; ds.
XX
OS Homo sapiens.
XX
PN WO200218632-A2.
XX
PD 07-MAR-2002.
XX
PF 01-SEP-2001; 2001WO-EP010074.
XX
PR 01-SEP-2000; 2000DE-01043826.
PR 05-SEP-2000; 2000DE-01044543.
XX
PA (EPIC-) EPIGENOMICS AG.
XX
PI Olek A, Piepenbrock C, Berlin K, Guetig D;
XX
DR WPI; 2002-371829/40.
XX
PT Determining the degree of cytosine methylation in genomic DNA, useful for
PT diagnosis and prognosis, comprises selective hybridization of amplicons
PT from chemically treated DNA.
XX
PS Claim 12; 56pp + Sequence Listing; 56pp; German.

CC This invention describes a novel method for determining the degree of
CC methylation of a particular cytosine in a motif 5'-CpG-3', present in a
CC genomic sample of DNA. The sample is treated chemically to convert
CC cytosine (C) but not methylated C, to uracil, then part of the genomic
CC DNA that contains the target C is amplified to form a labeled amplicon.
CC The amplicon is hybridised to two classes, each with at least one member,
CC of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers and the
CC degree of hybridisation to both classes is determined from the label on
CC the amplicon. From the ratio of labels hybridised to the two classes of
CC oligomers, the degree of methylation is calculated. The method is used:
CC (i) for diagnosis and/or prognosis of side effects of therapeutic drugs
CC and of a wide range of diseases, e.g. cancer, disorders of the central
CC nervous, cardiovascular, gastrointestinal and respiratory systems etc.,

CC particularly by detecting mutations or single nucleotide polymorphisms
CC (SNP's); and (ii) for differentiation of cell or tissue types and for
CC investigating cell differentiation. The method allows the methylation
CC status of many C residues to be determined simultaneously. ABO13410-
CC ABO54121 represent genomic DNA sequences used to illustrate the method
CC for determining the degree of cytosine methylation described in the
CC disclosure of the invention

XX Sequence 618 BP; 115 A; 58 C; 204 G; 241 T; 0 U; 0 Other;

Query Match Best Local Similarity 92.7%; Score 20.4; DB 6; Length 618;

Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TCCTCGTTTTTCGTGCGTTTTT 22
| | | | | | | | | | | | | | | | | | | | | |
Db TCCTCGTTTTTCGTGCGTTTTT 119

RESULT 6
ID ABO53697/c
XX ABO53697 standard; DNA; 619 BP.

XX ABO53697;

DT 12-JUL-2002 (first entry)

DE Oligonucleotide for detecting cytosine methylation SEQ ID NO 40288.

XX Human; cytosine methylation; 5'-CpG-3'; uracil; cytosine; diagnosis;
XX drug; side effect; cancer; central nervous system; cardiovascular;
XX gastrointestinal; respiratory system; single nucleotide polymorphism;
XX SNP; cell differentiation; ds.

OS Homo sapiens.

PN WO200218632-A2.

PD 07-MAR-2002.

PF 01-SEP-2001; 2001WO-EP010074.

PR 01-SEP-2000; 2000DE-01043826.

PR 05-SEP-2000; 2000DE-01044543.

PA (EPIG-) EPIGENOMICS AG.

PI Olek A, Piepenbrock C, Berlin K, Guefig D;

DR WPI; 2002-371829/40.

PT Determining the degree of cytosine methylation in genomic DNA, useful for
PT diagnosis and prognosis, comprises selective hybridization of amplicons
PT from chemically treated DNA.

PS Claim 12; 56pp + Sequence Listing; 56pp; German.

XX This invention describes a novel method for determining the degree of
CC methylation of a particular cytosine in a motif 5'-CpG-3', present in a
CC genomic sample of DNA. The sample is treated chemically to convert
CC cytosine (C) but not methylated C, to uracil, then part of the genomic
CC DNA that contains the target C is amplified to form a labeled amplicon.
CC The amplicon is hybridized to two classes, each with at least one member,
CC of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers and the
CC degree of hybridization to both classes is determined from the label on
CC the amplicon. From the ratio of labels hybridized to the two classes of
CC oligomers, the degree of methylation is calculated. The method is used:
CC (i) for diagnosis and/or prognosis of side effects of therapeutic drugs
CC and of a wide range of diseases, e.g. cancer, disorders of the central
CC nervous, cardiovascular, gastrointestinal and respiratory systems etc.,
CC particularly by detecting mutations or single nucleotide polymorphisms
CC (SNP's); and (ii) for differentiation of cell or tissue types and for
CC investigating cell differentiation. The method allows the methylation

CC status of many C residues to be determined simultaneously. ABO13410-
CC ABO54121 represent genomic DNA sequences used to illustrate the method
CC for determining the degree of cytosine methylation described in the
CC disclosure of the invention

XX Sequence 619 BP; 241 A; 205 C; 57 G; 116 T; 0 U; 0 Other;

Query Match Best Local Similarity 92.7%; Score 20.4; DB 6; Length 619;

Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TCCTCGTTTTTCGTGCGTTTTT 22
| | | | | | | | | | | | | | | | | | | | | |
Db 430 TCCTCGTTTTTCGTGCGTTTTT 409

RESULT 7
ID ABO53696
XX ABO53696 standard; DNA; 619 BP.

XX ABO53696;

DT 12-JUL-2002 (first entry)

DE Oligonucleotide for detecting cytosine methylation SEQ ID NO 40287.

XX Human; cytosine methylation; 5'-CpG-3'; uracil; cytosine; diagnosis;
XX drug; side effect; cancer; central nervous system; cardiovascular;
XX gastrointestinal; respiratory system; single nucleotide polymorphism;
XX SNP; cell differentiation; ds.

OS Homo sapiens.

PN WO200218632-A2.

PD 07-MAR-2002.

PF 01-SEP-2001; 2001WO-EP010074.

PR 01-SEP-2000; 2000DE-01043826.

PR 05-SEP-2000; 2000DE-01044543.

PA (EPIG-) EPIGENOMICS AG.

PI Olek A, Piepenbrock C, Berlin K, Guefig D;

DR WPI; 2002-371829/40.

PT Determining the degree of cytosine methylation in genomic DNA, useful for
PT diagnosis and prognosis, comprises selective hybridization of amplicons
PT from chemically treated DNA.

PS Claim 12; 56pp + Sequence Listing; 56pp; German.

XX This invention describes a novel method for determining the degree of
CC methylation of a particular cytosine in a motif 5'-CpG-3', present in a
CC genomic sample of DNA. The sample is treated chemically to convert
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CC DNA that contains the target C is amplified to form a labeled amplicon.
CC The amplicon is hybridized to two classes, each with at least one member,
CC of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers and the
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CC the amplicon. From the ratio of labels hybridized to the two classes of
CC oligomers, the degree of methylation is calculated. The method is used:
CC (i) for diagnosis and/or prognosis of side effects of therapeutic drugs
CC and of a wide range of diseases, e.g. cancer, disorders of the central
CC nervous, cardiovascular, gastrointestinal and respiratory systems etc.,
CC particularly by detecting mutations or single nucleotide polymorphisms
CC (SNP's); and (ii) for differentiation of cell or tissue types and for
CC investigating cell differentiation. The method allows the methylation
CC status of many C residues to be determined simultaneously. ABO13410-
CC ABO54121 represent genomic DNA sequences used to illustrate the method
CC for determining the degree of cytosine methylation described in the

CC disclosure of the invention
XX Sequence 619 BP; 116 A; 57 C; 205 G; 241 T; 0 U; 0 Other;
SQ Query Match 92.7%; Score 20.4; DB 6; Length 619;
Best Local Similarity 95.5%; Pred. No. 26;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
OY 1 TCGTCGTTTTTCGTCGTTTTT 22
Db 190 TCGTCGTTTTTCGTCGTTTTT 211

RESULT 8
ABQ15435/C
ID ABQ15435 standard; DNA; 761 BP.
XX
XX ABQ15435;
AC
XX
XX 12-JUN-2002 (first entry)
DT
XX
XX Oligonucleotide for detecting cytosine methylation SEQ ID NO 2026.
DE
XX Human; cytosine methylation; 5'-CpG-3'; uracil; cytosine; diagnosis;
KW drug; side effect; cancer; central nervous system; cardiovascular;
KW gastrointestinal; respiratory system; single nucleotide polymorphism;
KW SNP; cell differentiation; ds.
XX
XX Homo sapiens.
OS
XX
XX WO200218632-A2.
XX
XX 07-MAR-2002.
XX
XX 01-SEP-2001; 2001WO-EP010074.
XX
XX 01-SEP-2000; 2000DE-01043826.
XX
XX 05-SEP-2000; 2000DE-01044543.
XX
XX (EPIC-) EPIGENOMICS AG.
XX
XX Olek A, Piepenbrock C, Berlin K, Guectig D;
XX WPI; 2002-371829/40.
XX
XX Determining the degree of cytosine methylation in genomic DNA, useful for
PT diagnosis and prognosis, comprises selective hybridization of amplicons
PT from chemically treated DNA.
XX
XX
XX Claim 12; 56pp + Sequence Listing; 56pp; German.

This invention describes a novel method for determining the degree of
CC methylation of a particular cytosine in a motif 5'-CpG-3', present in a
CC genomic sample of DNA. The sample is treated chemically to convert
CC cytosine (C) but not methylated C, to uracil, then part of the genomic
CC DNA that contains the target C is amplified to form a labeled amplicon.
CC The amplicon is hybridised to two classes, each with at least one member,
CC of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers and the
CC degree of hybridisation to both classes is determined from the label on
CC the amplicon. From the ratio of labels hybridised to the two classes of
CC oligomers, the degree of methylation is calculated. The method is used:
CC (i) for diagnosis and/or prognosis of side effects of therapeutic drugs
CC and of a wide range of diseases, e.g. cancer, disorders of the central
CC nervous, cardiovascular, gastrointestinal and respiratory systems etc.,
CC particularly by detecting mutations or single nucleotide polymorphisms
CC (SNP's); and (ii) for differentiation of cell or tissue types and for
CC investigating cell differentiation. The method allows the methylation
CC status of many C residues to be determined simultaneously. ABQ13410-
CC ABQ14121 represent genomic DNA sequences used to illustrate the method
CC for determining the degree of cytosine methylation described in the
CC disclosure of the invention
XX
XX Sequence 761 BP; 238 A; 332 C; 100 G; 91 T; 0 U; 0 Other;

Query Match 92.7%; Score 20.4; DB 6; Length 761;
Best Local Similarity 95.5%; Pred. No. 26;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
OY 1 TCGTCGTTTTTCGTCGTTTTT 22
Db 164 TCGTCGTTTTTCGTCGTTTTT 143

RESULT 9
ABQ15434
ID ABQ15434 standard; DNA; 761 BP.
XX
XX ABQ15434;
AC
XX
XX 12-JUN-2002 (first entry)
DT
XX
XX Oligonucleotide for detecting cytosine methylation SEQ ID NO 2025.
DE
XX Human; cytosine methylation; 5'-CpG-3'; uracil; cytosine; diagnosis;
KW drug; side effect; cancer; central nervous system; cardiovascular;
KW gastrointestinal; respiratory system; single nucleotide polymorphism;
KW SNP; cell differentiation; ds.
XX
XX Homo sapiens.
OS
XX
XX WO200218632-A2.
XX
XX 07-MAR-2002.
XX
XX 01-SEP-2001; 2001WO-EP010074.
XX
XX 01-SEP-2000; 2000DE-01043826.
XX
XX 05-SEP-2000; 2000DE-01044543.
XX
XX (EPIC-) EPIGENOMICS AG.
XX
XX Olek A, Piepenbrock C, Berlin K, Guectig D;
XX WPI; 2002-371829/40.
XX
XX Determining the degree of cytosine methylation in genomic DNA, useful for
PT diagnosis and prognosis, comprises selective hybridization of amplicons
PT from chemically treated DNA.
XX
XX
XX Claim 12; 56pp + Sequence Listing; 56pp; German.

This invention describes a novel method for determining the degree of
CC methylation of a particular cytosine in a motif 5'-CpG-3', present in a
CC genomic sample of DNA. The sample is treated chemically to convert
CC cytosine (C) but not methylated C, to uracil, then part of the genomic
CC DNA that contains the target C is amplified to form a labeled amplicon.
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CC of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers and the
CC degree of hybridisation to both classes is determined from the label on
CC the amplicon. From the ratio of labels hybridised to the two classes of
CC oligomers, the degree of methylation is calculated. The method is used:
CC (i) for diagnosis and/or prognosis of side effects of therapeutic drugs
CC and of a wide range of diseases, e.g. cancer, disorders of the central
CC nervous, cardiovascular, gastrointestinal and respiratory systems etc.,
CC particularly by detecting mutations or single nucleotide polymorphisms
CC (SNP's); and (ii) for differentiation of cell or tissue types and for
CC investigating cell differentiation. The method allows the methylation
CC status of many C residues to be determined simultaneously. ABQ13410-
CC ABQ14121 represent genomic DNA sequences used to illustrate the method
CC for determining the degree of cytosine methylation described in the
CC disclosure of the invention
XX
XX Sequence 761 BP; 91 A; 100 C; 332 G; 238 T; 0 U; 0 Other;

Query Match 92.7%; Score 20.4; DB 6; Length 761;
Best Local Similarity 95.5%; Pred. No. 26;

Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TCGTCGTTTTTCGTCGTTTTT 22
 |||||
 DB 598 TCGTCGTTTTTCGTCGTTTTT 619

RESULT 10
 ABO20612
 ID ABO20612 standard; DNA; 1024 BP.
 XX ABO20612;

XX 12-JUL-2002 (first entry)

DE Oligonucleotide for detecting cytosine methylation SEQ ID NO 7203.

XX Human; cytosine methylation; 5'-CpG-3'; uracil; cytosine; diagnosis;
 KM drug; side effect; cancer; central nervous system; cardiovascular;
 KM gastrointestinal; respiratory system; single nucleotide polymorphism;
 XX SNP; cell differentiation; ds.

XX Homo sapiens.

XX WO200218632-A2.

XX 07-MAR-2002.

PF 01-SEP-2001; 2001WO-EP010074.

XX 01-SEP-2000; 2000DE-01043826.

PR 05-SEP-2000; 2000DE-01044543.

XX (EPIC-) EPIGENOMICS AG.

PI Olek A, Piepenbrock C, Berlin K, Guetig D;

XX WPI; 2002-371829/40.

PT Determining the degree of cytosine methylation in genomic DNA, useful for
 PT diagnosis and prognosis, comprises selective hybridization of amplicons
 PT from chemically treated DNA.

PS Claim 12; 56bp + Sequence Listing; 56bp; German.

XX This invention describes a novel method for determining the degree of
 CC methylation of a particular cytosine in a motif 5'-CpG-3', present in a
 CC genomic sample of DNA. The sample is treated chemically to convert
 CC cytosine (C) but not methylated C, to uracil, then part of the genomic
 CC DNA that contains the target C is amplified to form a labeled amplicon.
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 CC the amplicon. From the ratio of labels hybridised to the two classes of
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 CC (i) for diagnosis and/or prognosis of side effects of therapeutic drugs
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 CC particularly by detecting mutations or single nucleotide polymorphisms
 CC (SNP's); and (ii) for differentiation of cell or tissue types and for
 CC investigating cell differentiation. The method allows the methylation
 CC status of many C residues to be determined simultaneously. ABO13410-
 CC ABO54121 represent genomic DNA sequences used to illustrate the method
 CC for determining the degree of cytosine methylation described in the
 CC disclosure of the invention

XX Sequence 1024 BP; 138 A; 122 C; 396 G; 368 T; 0 U; 0 Other;

Query Match 92.7%; Score 20.4; DB 6; Length 1024;

Best Local Similarity 95.5%; Pred. No. 26; Mismatches 1; Indels 0; Gaps 0;

Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TCGTCGTTTTTCGTCGTTTTT 22

DB 276 TCGTCGTTTTTCGTCGTTTTT 297
 |||||

RESULT 11
 ABO20613/C
 ID ABO20613 standard; DNA; 1024 BP.
 XX ABO20613;

XX 12-JUL-2002 (first entry)

DE Oligonucleotide for detecting cytosine methylation SEQ ID NO 7204.

XX Human; cytosine methylation; 5'-CpG-3'; uracil; cytosine; diagnosis;
 KM drug; side effect; cancer; central nervous system; cardiovascular;
 KM gastrointestinal; respiratory system; single nucleotide polymorphism;
 XX SNP; cell differentiation; ds.

XX Homo sapiens.

XX WO200218632-A2.

XX 07-MAR-2002.

PF 01-SEP-2001; 2001WO-EP010074.

XX 01-SEP-2000; 2000DE-01043826.

PR 05-SEP-2000; 2000DE-01044543.

XX (EPIC-) EPIGENOMICS AG.

PI Olek A, Piepenbrock C, Berlin K, Guetig D;

XX WPI; 2002-371829/40.

PT Determining the degree of cytosine methylation in genomic DNA, useful for
 PT diagnosis and prognosis, comprises selective hybridization of amplicons
 PT from chemically treated DNA.

PS Claim 12; 56bp + Sequence Listing; 56bp; German.

XX This invention describes a novel method for determining the degree of
 CC methylation of a particular cytosine in a motif 5'-CpG-3', present in a
 CC genomic sample of DNA. The sample is treated chemically to convert
 CC cytosine (C) but not methylated C, to uracil, then part of the genomic
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 CC the amplicon. From the ratio of labels hybridised to the two classes of
 CC oligomers, the degree of methylation is calculated. The method is used:
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 CC and of a wide range of diseases, e.g. cancer, disorders of the central
 CC nervous, cardiovascular, gastrointestinal and respiratory systems etc.,
 CC particularly by detecting mutations or single nucleotide polymorphisms
 CC (SNP's); and (ii) for differentiation of cell or tissue types and for
 CC investigating cell differentiation. The method allows the methylation
 CC status of many C residues to be determined simultaneously. ABO13410-
 CC ABO54121 represent genomic DNA sequences used to illustrate the method
 CC for determining the degree of cytosine methylation described in the
 CC disclosure of the invention

XX Sequence 1024 BP; 368 A; 396 C; 122 G; 138 T; 0 U; 0 Other;

Query Match 92.7%; Score 20.4; DB 6; Length 1024;

Best Local Similarity 95.5%; Pred. No. 26; Mismatches 1; Indels 0; Gaps 0;

Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TCGTCGTTTTTCGTCGTTTTT 22

DB 749 TCGTCGTTTTTCGTCGTTTTT 728

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RESULT 12
ABQ41425/C
ID ABQ41425 standard; DNA; 511 BP.
XX
XX ABQ41425;
AC
XX 12-JUL-2002 (first entry)
DT
XX Oligonucleotide for detecting cytosine methylation SEQ ID NO 28016.
DE
XX Human; cytosine methylation; 5'-CpG-3'; uracil; cytosine; diagnosis;
XX drug; side effect; cancer; central nervous system; cardiovascular;
XX gastrointestinal; respiratory system; single nucleotide polymorphism;
XX SNP; cell differentiation; ds.
XX
XX Homo sapiens.
OS
XX
XX WO200218632-A2.
XX
XX 07-MAR-2002.
XX
XX 01-SEP-2001; 2001WO-EP010074.
XX
XX 01-SEP-2000; 2000DE-01043826.
XX
XX 05-SEP-2000; 2000DE-01044543.
XX
XX (EPIG-) EPIGENOMICS AG.
XX
XX Olek A, Piepenbrock C, Berlin K, Guetig D;
XX
XX WPI; 2002-371829/40.
XX
XX Determining the degree of cytosine methylation in genomic DNA, useful for
XX diagnosis and prognosis, comprises selective hybridization of amplicons
XX from chemically treated DNA.
XX
XX Claim 12; 56pp + Sequence Listing; 56pp; German.
XX
XX This invention describes a novel method for determining the degree of
XX methylation of a particular cytosine in a motif 5'-CpG-3', present in a
XX genomic sample of DNA. The sample is treated chemically to convert
XX cytosine (C) but not methylated C, to uracil, then part of the genomic
XX DNA that contains the target C is amplified to form a labeled amplicon.
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XX degree of hybridisation to both classes is determined from the label on
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XX oligomers, the degree of methylation is calculated. The method is used:
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XX nervous, cardiovascular, gastrointestinal and respiratory systems etc.,
XX particularly by detecting mutations or single nucleotide polymorphisms
XX (SNP's); and (ii) for differentiation of cell or tissue types and for
XX investigating cell differentiation. The method allows the methylation
XX status of many C residues to be determined simultaneously. ABQ13410-
XX ABQ41421 represent genomic DNA sequences used to illustrate the method
XX for determining the degree of cytosine methylation described in the
XX disclosure of the invention
XX
XX Sequence 511 BP; 180 A; 177 C; 62 G; 92 T; 0 U; 0 Other;
XX
XX Query Match 88.2%; Score 19.4; DB 6; Length 511;
XX Best Local Similarity 95.2%; Pred. No. 70;
XX Matches 20; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
```

```
ID ABQ41424 standard; DNA; 511 BP.
XX
XX ABQ41424;
AC
XX 12-JUL-2002 (first entry)
DT
XX Oligonucleotide for detecting cytosine methylation SEQ ID NO 28015.
DE
XX Human; cytosine methylation; 5'-CpG-3'; uracil; cytosine; diagnosis;
XX drug; side effect; cancer; central nervous system; cardiovascular;
XX gastrointestinal; respiratory system; single nucleotide polymorphism;
XX SNP; cell differentiation; ds.
XX
XX Homo sapiens.
OS
XX
XX WO200218632-A2.
XX
XX 07-MAR-2002.
XX
XX 01-SEP-2001; 2001WO-EP010074.
XX
XX 01-SEP-2000; 2000DE-01043826.
XX
XX 05-SEP-2000; 2000DE-01044543.
XX
XX (EPIG-) EPIGENOMICS AG.
XX
XX Olek A, Piepenbrock C, Berlin K, Guetig D;
XX
XX WPI; 2002-371829/40.
XX
XX Determining the degree of cytosine methylation in genomic DNA, useful for
XX diagnosis and prognosis, comprises selective hybridization of amplicons
XX from chemically treated DNA.
XX
XX Claim 12; 56pp + Sequence Listing; 56pp; German.
XX
XX This invention describes a novel method for determining the degree of
XX methylation of a particular cytosine in a motif 5'-CpG-3', present in a
XX genomic sample of DNA. The sample is treated chemically to convert
XX cytosine (C) but not methylated C, to uracil, then part of the genomic
XX DNA that contains the target C is amplified to form a labeled amplicon.
XX The amplicon is hybridised to two classes, each with at least one member,
XX of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers and the
XX degree of hybridisation to both classes is determined from the label on
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XX (i) for diagnosis and/or prognosis of side effects of therapeutic drugs
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XX particularly by detecting mutations or single nucleotide polymorphisms
XX (SNP's); and (ii) for differentiation of cell or tissue types and for
XX investigating cell differentiation. The method allows the methylation
XX status of many C residues to be determined simultaneously. ABQ13410-
XX ABQ41421 represent genomic DNA sequences used to illustrate the method
XX for determining the degree of cytosine methylation described in the
XX disclosure of the invention
XX
XX Sequence 511 BP; 92 A; 62 C; 177 G; 180 T; 0 U; 0 Other;
XX
XX Query Match 88.2%; Score 19.4; DB 6; Length 511;
XX Best Local Similarity 95.2%; Pred. No. 70;
XX Matches 20; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
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RESULT 14
ABQ48469/C
ID ABQ48469 standard; DNA; 523 BP.
XX
XX ABQ48469;
AC
```

[illegible]

DE Oligonucleotide for detecting cytosine methylation SEQ ID NO 35059.
XX Human; cytosine methylation; 5'-CpG-3'; uracil; cytosine; diagnosis;
XX drug; side effect; cancer; central nervous system; cardiovascular;
XX gastrointestinal; respiratory system; single nucleotide polymorphism;
XX SNP; cell differentiation; ds.
OS Homo sapiens.
XX
XX WO200218632-A2.
XX
XX 07-MAR-2002.
XX
XX 01-SEP-2001; 2001WO-EP010074.
XX
XX 01-SEP-2000; 2000DE-01043826.
XX 05-SEP-2000; 2000DE-01044543.
XX
XX (EP1G-) EPIGENOMICS AG.
XX
XX Olek A, Piepenbrock C, Berlin K, Guetig D;
XX WPI; 2002-371829/40.
XX
XX Determining the degree of cytosine methylation in genomic DNA, useful for
XX diagnosis and prognosis, comprises selective hybridization of amplicons
XX from chemically treated DNA.
XX
XX Claim 12; 56pp + Sequence Listing; 56pp; German.
XX
XX This invention describes a novel method for determining the degree of
XX methylation of a particular cytosine in a motif 5'-CpG-3', present in a
XX genomic sample of DNA. The sample is treated chemically to convert
XX cytosine (C) but not methylated C, to uracil, then part of the genomic
XX DNA that contains the target C is amplified to form a labeled amplicon.
XX The amplicon is hybridized to two classes, each with at least one member,
XX of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers and the
XX degree of hybridization to both classes is determined from the label on
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XX oligomers, the degree of methylation is calculated. The method is used:
XX (i) for diagnosis and/or prognosis of side effects of therapeutic drugs
XX and of a wide range of diseases, e.g. cancer, disorders of the central
XX nervous, cardiovascular, gastrointestinal and respiratory systems etc.,
XX particularly by detecting mutations or single nucleotide polymorphisms
XX (SNP's); and (ii) for differentiation of cell or tissue types and for
XX investigating cell differentiation. The method allows the methylation
XX status of many C residues to be determined simultaneously. AB013410-
XX AB05411 represent genomic DNA sequences used to illustrate the method
XX for determining the degree of cytosine methylation described in the
XX disclosure of the invention
XX
XX Sequence 523 BP; 64 A; 61 C; 207 G; 191 T; 0 U; 0 Other;
XX
XX Query Match 88.2%; Score 19.4; DB 6; Length 523;
XX Best Local Similarity 95.2%; Pred. No. 70;
XX Matches 20; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
XX
XX 2 CGTCGTTTTTCGTCGTTTTT 22
XX ||||||| |||||||
XX Db 488 CGTCGTTTTTCGTCGTTTTT 508
XX

Search completed: March 9, 2005, 17:40:33
Job time : 537 secs